

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:06:39 ; Search time 15 Seconds
(without alignments)
1769.504 Million cell updates/sec

Title: US-10-092-404-2
Perfect score: 1520
Sequence: 1 RLRSLSLHYLFMGASEQDL.....RYTCQVEHPGLDPLIVWE 276

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 76:.*
1: pir1:.*
2: pir2:.*
3: pir3:.*
4: pir4:.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	ID	Description
1	1140	75.0	359	2 JCS382	hereditary hemochromatosis protein precursor - mouse
2	542.5	35.7	341	2 A57136	class I histocompa
3	523	34.4	361	1 HLRB	MHC class I histoc
4	523	34.4	361	2 I46858	MHC class I RLA pr
5	520	34.2	332	2 S06424	MHC class I histoc
6	517	34.0	365	2 I36961	MHC class I protei
7	516	33.9	361	2 B27638	MHC class I histoc
8	515	33.9	365	2 I83063	A11.2 - human
9	514	33.8	365	2 A47636	MHC class I histoc
10	514	33.8	365	2 I56039	HLA-A30.3 precurs
11	512	33.7	370	1 HLHUA3	MHC class I histoc
12	510	33.6	365	2 I38439	MHC class I histoc
13	509	33.5	365	2 I37542	MHC class I histoc
14	509	33.5	365	2 I38442	gene HLA-A-0205 pr
15	509	33.5	365	2 I61902	MHC class I histoc
16	508	33.4	365	2 I72170	MHC class I histoc
17	508	33.4	365	2 I38441	gene HLA-A-6802 pr
18	506	33.3	365	1 HLHUA2	MHC class I histoc
19	506	33.3	365	2 I37482	MHC class I histoc
20	506	33.3	365	2 I38519	MHC class I histoc
21	506	33.3	365	2 I84448	MHC class I histoc
22	505	33.2	365	2 I38610	MHC class I histoc
23	505	33.2	365	2 I37470	HLA-A*0210 - human
24	504	33.2	355	2 T28149	MHC class I histoc
25	504	33.2	364	2 S03535	class I histocompa
26	503	33.1	365	2 I37476	MHC class I histoc
27	503	33.1	365	2 I37478	MHC class I histoc
28	503	33.1	365	2 I38443	gene HLA-A-0203 pr
29	503	33.1	365	2 I61857	MHC HLA-A2.4a chai

30	502.5	33.1	341	2 JCS663	major histocompati
31	502	33.0	357	2 I36965	MHC class I protei
32	501.5	33.0	362	2 A45845	MHC class I histoc
33	501	33.0	365	2 I61856	MHC class I histoc
34	501	33.0	365	2 I54493	MHC class I histoc
35	500	32.9	273	1 HLHUE9	MHC class I histoc
36	500	32.9	365	2 S77963	MHC class I histoc
37	500	32.9	365	2 S01171	class I histocompa
38	500	32.9	365	2 I54416	HLA-AW34 protein -
39	499	32.8	365	2 I37483	HLA-AW34.2 antigen
40	498	32.8	273	1 HLHUAW	MHC class I histoc
41	498	32.8	360	2 A27638	MHC class I histoc
42	498	32.8	365	2 I72171	HLA-AW33.1, HLA-Aw
43	497.5	32.7	339	2 I56071	MHC class I histoc
44	497	32.7	279	2 JX0353	zinc-alpha 2-glyco
45	497	32.7	362	2 I68724	MHC class I histoc

ALIGNMENTS

RESULT 1

JCS382
hereditary hemochromatosis protein precursor - mouse
C:Species: Mus musculus (house mouse)
C>Date: 02-Jun-1997 #sequence_revision 18-Jul-1997 #text_change 05-Nov-1999
C:Accession: JCS382
R:Hashimoto, K.; Hirai, M.; Kurosawa, Y.
Biochem. Biophys. Res. Commun. 230, 35-39, 1997
A:Title: Identification of a mouse homolog for the human hereditary haemochromatosis ca
A:Reference number: JCS382; MUID:97148566; PMID:9020055
A:Accession: JCS382
A:Status: nucleic acid sequence not shown
A:Molecule type: DNA
A:Residues: 1-359 <HAS>
A:Cross-references: GB:U66849; NID:g1519484; PIDN:AAB07525.1; PID:g1519485
C:Comment: This protein plays a role in iron metabolism.
C:Genetics:
A:Gene: mr2
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
F:1-29/Domain: signal sequence #status predicted <SIG>
F:30-359/Product: hereditary haemochromatosis protein #status predicted <MAT>
F:30-117/Domain: alpha 1 #status predicted <ALF1>
F:118-217/Domain: alpha 2 #status predicted <ALF2>
F:218-309/Domain: alpha 3 #status predicted <ALF3>
F:314-340/Domain: transmembrane #status predicted <TRM>
F:341-359/Domain: intracellular #status predicted <INT>

Query Match 75.0%; Score 1140; DB 2; Length 359;

Best Local Similarity 72.2%; Pred. No. 2.3e-87;
Matches 203; Conservative 30; Mismatches 40; Indels 8; Gaps 1;

QY	4	RSLSLHYLFMGASEODLGLSFEALGYDDQLFVYDDSRVRVPTWVSRISSQWL	63
DB	30	RSLSLHYLFMGASEPDGLPLFEARGYDDQLFVSYNHSRAEPRAWILEQTSQWL	89
QY	64	QLSQSLKGDHMTFTVDFWTIMENHNSK-----ESHTLQVILGCEQDNSTEGYWK	115
DB	90	HLSQLKGDYMFVDFWTIMGNVNSKVTGLGVVSESHILQVILGCEVHEDNSTSGFWR	149
QY	116	GYDQGDHLFCFDPDILDRAPRAWPTKLEWERIKIRARQNRYLERDCPAQLQLLEL	175
DB	150	GYDQGDHLFCFCKTLNWSAEPGAWATKVEWDEHKIRAKQNRDYLEKDCPQLRLLEL	209
QY	176	GRGVLDQVPLVVKVTHVTSSVTLRCALNYPQNTIMKWLKDKQPMDAKEFPKDV	235
DB	210	GRGVLDQVPTLVKTRHWTSGTSLRQALDFFPQNTIMRWLKNQNPDLKDVNPEKVL	269
QY	236	PNGDGTQGWITLAVPPGEEQRYTCQVEHPGLDQLPLIWI	276
DB	270	PNGDGTQGWITLAVAPGDETRFTCQVEHPGLDQLPLTASWE	310

RESULT 2

A57136
Class I histocompatibility antigen related protein MR1 precursor - human
C:Species: Homo sapiens (man)
C:Date: 23-Feb-1996 #sequence_revision 23-Feb-1996 #text_change 23-Jul-1999
C:Accession: A57136
R:Hashimoto, K.; Hirai, M.; Kurosawa, Y.
Science 269, 693-695, 1995
A:Title: A gene outside the human MHC related to classical HLA class I genes.
A:Reference number: A57136; MUID:95350662; PMID:7624800
A:Accession: A57136
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-341 <HAS>
A:Cross-references: GB:U22963; NID:9940353; PIDN:AA50174.1; PID:g940354
C:Genetics:
A:Gene: GDB:HLALS
A:Cross-references: GDB:683188; OMIM:600764
A:Map position: 1q25.3-1q25.3
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 35.78; Score 542.5; DB 2; Length 341;
Best Local Similarity 39.5%; Pred. No. 1.2e-37;
Matches 107; Conservative 50; Mismatches 111; Indels 3; Gaps 3;

QY 4 RSHSLYLFMGASEQDGLSLFEALGVDDQLFVYDDESRVPEPTPWSSRISSQMWL 63
DB 23 RTHSLRFLGVSDPHGVPEFISGVYDSDHPITTYDSVTRQKPEAPWNAENLADHWE 82
QY 64 QLSQSLKGDHMFVDFWITMENHNSKE-SHTLQVILGCEMQEDNSTEGYWKYGYDGDQ 123
DB 83 RYTQLRGWQMFVKELKRLQRHNS-GSHTYQRMIGCELLEDGSTTGLQYAYDQDP 141
QY 124 LEPCPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDCAQQLLELGRGVLDQ 183
DB 142 LIENKDTLSLWADVNAHTIKQAEANQHLLYQKNLEBECIAWLKRFLEYKDTLQRT 201
QY 184 VPPLVAVKTHVT-SSVTLRCALNYYPQNTWKWLKQKPMDAKEFEKPDVLPNGDGY 242
DB 202 EPPLVVRNRKETFPFGVTFALCKAHGFYPPEIYMTWMKNGEEI-VOEIDYGDILPVG 260
QY 243 QGMWITLAVPGEQRYTCQVEHPGLDQPLIV 273
DB 261 QAWASIELDQSSNLNLYSCHVEHCGVHMLQV 291

RESULT 3

HLRB
MHC class I histocompatibility antigen RLA alpha chain precursor (RL-5) - rabbit
C:Species: Oryctolagus cuniculus (domestic rabbit)
C:Date: 25-Feb-1985 #sequence_revision 25-Feb-1985 #text_change 22-Jun-1999
C:Accession: A02193
R:Tykocinski, M.L.; Marche, P.N.; Max, E.E.; Kindt, T.J.
J. Immunol. 133, 2261-2269, 1984
A:Title: Rabbit class I MHC genes: cDNA clones define full-length transcripts of an expressed polymorphic gene.
A:Reference number: A02193; MUID:84290724; PMID:6432910
A:Accession: A02193
A:Molecule type: mRNA
A:Residues: 1-361 <TYK>
A:Cross-references: GB:K02441; NID:g1293894; PIDN:AAA98729.1; PID:g165496
A:Note: The source of this protein is a T-lymphoid cell line (RL-5), which has been transferred to a new cell line (RL-5) and humanized.
C:Comment: In contrast to the many antigens expressed in mouse (K, D, and L) and human (A, B, and C) class I MHC molecules, the RLA alpha chain has a unique structure. The RLA alpha chain may therefore differ from the HLA and H-2 loci in having limited complexity.
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
C:Keywords: duplication; glycoprotein; heterodimer; transmembrane protein
F:1-24/Domain: signal sequence #status predicted <SIG>
F:25-361/Product: class I histocompatibility antigen RLA alpha chain #status predicted <EXT>
F:25-307/Domain: extracellular #status predicted <EXT>
F:25-114/Domain: alpha-1 <EX1>
F:115-206/Domain: alpha-2 <EX2>
F:220-285/Domain: immunoglobulin homology <IMM>
F:308-329/Domain: transmembrane #status predicted <TM>
F:330-361/Domain: intracellular #status predicted <INT>

F:110/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:125-188,227-283/Disulfide bonds: #status predicted

Query Match 34.4%; Score 523; DB 1; Length 361;
Best Local Similarity 40.1%; Pred. No. 5.3e-36;
Matches 111; Conservative 44; Mismatches 114; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDGLSLFEALGVDDQLFVYDDE--SRVPEPTPWSSRISSQMW 62
DB 26 SHSMRYFTYSVRPGLEPRFIIVGVYDVTQVFRDSDAASPRMEQAPWM-QQVEPEY 84
QY 63 LQLSLSKLGWDMHMFVDFWITMENHNSKE-SHTLQVILGCEMQEDNSTEGYWKYGYD 120
DB 85 DQQTQIAKDTAQTFRVNLNTALRYYNQSAAGSHTTQTMFGCEVWADGRFFHGYRQYAYD 144
QY 121 QHLEFPCPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDCAQQLLELGRGV 180
DB 145 ADYIALNEDLRSWTAADTAQNTQKWEAAG-EAERHAYLRECEVWLRRLYLEMGKETL 203
QY 181 DQOVPEPLVKVTHHTVSS-VTTLRCALNYYPQNTWKWLKQKPMDAKEFEKPDVLPNGD 239
DB 204 QRADPPKAVTHHPASDREATLRCWALGFYPAEISLTWQDGED-QTQDTLVELVTRPGD 262
QY 240 GTYQGWITLAVPGEQRYTCQVEHPGLDQPLIVWE 276
DB 263 GTFOKAAVVPVSGEQRVTCRVQHEGLPEPLTLTWE 299

RESULT 4

I46858
MHC class I RLA precursor - rabbit
C:Species: Oryctolagus cuniculus (domestic rabbit)
C:Date: 14-Feb-1997 #sequence_revision 14-Feb-1997 #text_change 21-Jan-2000
C:Accession: I46858
R:Marche, P.N.; Tykocinski, M.L.; Max, E.E.; Kindt, T.J.
Immunogenetics 21, 71-82, 1985
A:Title: Structure of a functional rabbit class I MHC gene: Similarity to human class I A;Reference number: I46858; MUID:85103547; PMID:3917574
A:Accession: I46858
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-361 <MAR>
A:Cross-references: GB:K02819; NID:g165497; PIDN:AAA98730.1; PID:g165498
C:Genetics:
A:Introns: 25/1; 115/1; 207/1; 299/1; 337/1; 348/1
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
F:220-285/Domain: immunoglobulin homology <IMM>

Query Match 34.4%; Score 523; DB 2; Length 361;
Best Local Similarity 40.1%; Pred. No. 5.3e-36;
Matches 111; Conservative 44; Mismatches 114; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDGLSLFEALGVDDQLFVYDDE--SRVPEPTPWSSRISSQMW 62
DB 26 SHSMRYFTYSVRPGLEPRFIIVGVYDVTQVFRDSDAASPRMEQAPWM-QQVEPEY 84
QY 63 LQLSLSKLGWDMHMFVDFWITMENHNSKE-SHTLQVILGCEMQEDNSTEGYWKYGYD 120
DB 85 DQQTQIAKDTAQTFRVNLNTALRYYNQSAAGSHTTQTMFGCEVWADGRFFHGYRQYAYD 144
QY 121 QHLEFPCPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDCAQQLLELGRGV 180
DB 145 ADYIALNEDLRSWTAADTAQNTQKWEAAG-EAERHAYLRECEVWLRRLYLEMGKETL 203
QY 181 DQOVPEPLVKVTHHTVSS-VTTLRCALNYYPQNTWKWLKQKPMDAKEFEKPDVLPNGD 239
DB 204 QRADPPKAVTHHPASDREATLRCWALGFYPAEISLTWQDGED-QTQDTLVELVTRPGD 262
QY 240 GTYQGWITLAVPGEQRYTCQVEHPGLDQPLIVWE 276
DB 263 GTFOKAAVVPVSGEQRVTCRVQHEGLPEPLTLTWE 299

RESULT 5

S06424
MHC class I histocompatibility antigen Ch25 alpha chain precursor - chimpanzee
N;Alternate names: MHC Ch1A chain
C;Species: Pan troglodytes (chimpanzee)
C;Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 23-Jul-1999
C;Accession: S06424; I36959
R;Lawlor, D.A.; Ward, F.E.; Ennis, P.D.; Jackson, A.P.; Parham, P.
Nature 335, 268-271, 1988
A;Title: HLA-A and B polymorphisms predate the divergence of humans and chimpanzees.
A;Reference number: S06424; MUID:98319000; PMID:3412487

A;Molecule type: mRNA
A;Residues: 1-332 <LAW>
R;Parham, P.; Lawlor, D.A.; Lomen, C.E.; Ennis, P.D.
J. Immunol. 142, 3937-3950, 1989
A;Title: Diversity and diversification of HLA-A,B,C alleles.
A;Reference number: I36956; MUID:99235215; PMID:2715640
A;Accession: I36959
A;Molecule type: mRNA
A;Residues: 1-332 <RES>
A;Cross-references: GB:M24047; NID:g176818; PID:AAA35426.1; PID:g553155
C;Superfamily: class I histocompatibility antigen; immunoglobulin homology
C;Keywords: glycoprotein; membrane protein
F;1-24/Domain: signal sequence #status predicted <SIG>
F;25-114/Domain: alpha-1 #status predicted <EX1>
F;115-206/Domain: alpha-2 #status predicted <EX2>
F;220-285/Domain: immunoglobulin homology <IMM>
F;307-331/Domain: transmembrane #status predicted <TM>
F;110/Binding site: carbohydrate (Asn) (covalent) #status predicted
F;125-188,227-283/Disulfide bonds: #status predicted

Query Match 34.2%; Score 520; DB 2; Length 332;
Best Local Similarity 40.1%; Pred. No. 8.5e-36;
Matches 111; Conservative 44; Mismatches 114; Indels 8; Gaps 7;

QY 5 SLSHLVFMGASQDGLSLFEALGYVDDQLFVYDDE--SRVPRTPWSSRISQMW 62
DB 26 SHSMRYFFTSVSRPGRGEPRFIAVGYYDDTQFVRFDSDAASQRMPEAPWIEQ-GPEYW 84
QY 63 LQLSLSLKGWDHMTVDFTWIMENHNHNSKE-SHTLQVILGCMEQDENS-TEGYWKYGDG 120
DB 85 DQETRAKAHSQTRDVLGTLRGYINQSGSHTIQIMYCDVGSQGRFLRGYRQDAYDG 144
QY 121 QDLHLEPCPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 KDVIALLNEDLRSWTAADMAAQITKRWEAAH-AAEQRAYLEGTVCVWELRRYLENGKETL 203
QY 181 DQVPPPLVKVTHH-VTSSVTTLCRAINYPPQITMKWLKDKQPMDAKEPEPKDVLNPGD 239
DB 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTELVELTRPAGD 262
QY 240 GTYQGMITLAVPGEORVTCVVEHGLDPLIVINE 276
DB 263 GTFQKAAVVVPSGEGEORYTCHVQHEGLPKPLTLRWE 299

RESULT 6

I36961
MHC class I protein - chimpanzee
C;Species: Pan troglodytes (chimpanzee)
C;Date: 04-Oct-1996 #sequence_revision 04-Oct-1996 #text_change 21-Jan-2000
C;Accession: I36961
R;Lawlor, D.A.; Warren, E.; Ward, F.E.; Parham, P.
Immunol. Rev. 113, 147-185, 1990
A;Title: Comparison of class I MHC alleles in humans and apes.
A;Reference number: I36961; MUID:90201944; PMID:1690682
A;Accession: I36961
A;Status: preliminary; translated from GB/EMBL/DBJ
A;Molecule type: mRNA
A;Residues: 1-365 <RES>
A;Cross-references: GB:M30678; NID:g176822; PID:AAA87970.1; PID:g176823
C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 34.0%; Score 517; DB 2; Length 365;
Best Local Similarity 39.7%; Pred. No. 1.7e-35;
Matches 110; Conservative 45; Mismatches 114; Indels 8; Gaps 7;
QY 5 SLSHLVFMGASQDGLSLFEALGYVDDQLFVYDDE--SRVPRTPWSSRISQMW 62
DB 26 SHSMRYFFTSVSRPGRGEPRFIAVGYYDDTQFVRFDSDAASQRMPEAPWIEQ-GPEYW 84
QY 63 LQLSLSLKGWDHMTVDFTWIMENHNHNSKE-SHTLQVILGCMEQDENS-TEGYWKYGDG 120
DB 85 DEETRAKAHSQTRDVLGTLRGYINQSGSHTIQIMYCDVGSQGRFLRGYRQDAYDG 144
QY 121 QDLHLEPCPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 KDVIALLNEDLRSWTAADMAAQITKRWEAAH-AAEQRAYLEGTVCVWELRRYLENGKETL 203
QY 181 DQVPPPLVKVTHH-VTSSVTTLCRAINYPPQITMKWLKDKQPMDAKEPEPKDVLNPGD 239
DB 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTELVELTRPAGD 262
QY 240 GTYQGMITLAVPGEORVTCVVEHGLDPLIVINE 276
DB 263 GTFQKAAVVVPSGEGEORYTCHVQHEGLPKPLTLRWE 299

RESULT 7

B27638
MHC class I histocompatibility antigen alpha chain precursor (BL3-7) - bovine
C;Species: Bos primigenius taurus (cattle)
C;Date: 08-Mar-1989 #sequence_revision 08-Mar-1989 #text_change 16-Feb-1997
C;Accession: B27638
R;Ennis, P.D.; Jackson, A.P.; Parham, P.
J. Immunol. 141, 642-651, 1988
A;Title: Molecular cloning of bovine class I MHC cDNA.
A;Reference number: A92826; MUID:88258075; PMID:3133413
A;Accession: B27638
A;Status: not compared with conceptual translation
A;Molecule type: mRNA
A;Residues: 1-361 <ENN>
C;Superfamily: class I histocompatibility antigen; immunoglobulin homology
C;Keywords: heterodimer; transmembrane protein
F;1-24/Domain: signal sequence #status predicted <SIG>
F;25-361/Product: MHC class I histocompatibility antigen, BOLA alpha chain (BL3-7) #status predicted
F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.9%; Score 516; DB 2; Length 361;
Best Local Similarity 38.9%; Pred. No. 2e-35;
Matches 109; Conservative 50; Mismatches 113; Indels 8; Gaps 7;

QY 2 LLRSLSHLVFMGASQDGLSLFEALGYVDDQLFVYDDE--SRVPRTPWSSRIS 59
DB 23 LAGSHSLRYFFTVGSVSRPGRGEPRFIAVGYYDDTQFVRFDSDAPNPREPRVPMWEGE-GP 81
QY 60 QMWLQLSLSLKGWDHMTVDFTWIMENHNHNSKE-SHTLQVILGCMEQDENS-TEGYWKY 117
DB 82 EYWRDNTRIYKDTAQIFRVDLNTLRGYINQSGSHNIQAMYGCDVGPGRLLRGFWQFG 141
QY 118 YDQDHLPECPPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGR 177
DB 142 YDGRDYIALNEELRSWTAADTAATQITKRWEAAH-AAETWRNYLEGECVWELRRYLENGK 200
QY 178 GYLDQVQVPLVKVTHH-VTSSVTTLCRAINYPPQITMKWLKDKQPMDAKEPEPKDVLN 236
DB 201 DTLRADPPKAHVTHHSISDREVTLCWALGFYPPEISLTWQREGD-QTQDMELVELTRP 259
QY 237 NGDGTVQGMITLAVPGEORVTCVVEHGLDPLIVINE 276
DB 260 SDGTFQKAAVVVPSGEGEORYTCHVQHEGLPKPLTLRWE 299

RESULT 8

I83063

All.2 - human
 C:Species: Homo sapiens (man)
 C>Date: 02-Aug-1996 #sequence_revision 02-Aug-1996 #text_change 21-Jan-2000
 C:Accession: I83063
 R:Lin, L.; Tokunaga, K.; Ishikawa, Y.; Bannai, M.; Kashiwase, K.; Kuwata, S.; Akaza, T.;
 Tissue Antigens 43, 78-82, 1994
 A>Title: Sequence analysis of serological HLA-A11 split antigens, All.1 and All.2.
 A:Reference number: I60129; MUID:94287401; PMID:8016845

A:Accession: I83063
 A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: mRNA
 A:Residues: 1-365 <RES>

A:Cross-references: GB:D16842; NID:G540517; PIDN:BA04118.1; PID:G487911
 C:Genetics:

A:Gene: A1102
 A:Superfamily: class I histocompatibility antigen; immunoglobulin homology
 F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.9%; Score 515; DB 2; Length 365;
 Best Local Similarity 39.4%; Pred. No. 2.5e-35;
 Matches 109; Conservative 47; Mismatches 113; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDE--SRVPERTPWVSSRISSQW 62
 DB 26 SHSMRYFYTSVRGCRKPRFIAVGYDDTQVRFDSDAASQRMPEAPWIEQE-GPEYW 84

QY 63 LQLSLSLQKGDHMFVDFWTIMENHNHKE-SHTLQVILGCEMQEDNS-TEGWYKYGYDG 120
 DB 85 DQETRNVAQSDTRVDLGLTRGYNQSDGSHTIQIMYGCDDVPGDGRFLGRQDAYDG 144

QY 121 QHLEFCPDTLDWRAAPRAWPKLEWERHKIRARONRAYLERDCPAQLQQLLELGRVL 180
 DB 145 KDYALNEDLRSWTAADMAAQITKRWAAH-AAEQRAYLEGRVLEWLRYLENGKETL 203

QY 181 DQVPPPLVKVTHH-VTSSVTLRCALNYYPNITMKWLKDKQPMDAKEPEPKDVLNPD 239
 DB 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTVELVETRPAGD 262

QY 240 GTYQGWITLAVPPGEORYTCQVHPGLDPLIVWE 276
 DB 263 GTFQKAAVVVPSGGEORYTCHVQHEGLPKPLTLRWE 299

RESULT 9
 A47636
 MHC class I histocompatibility antigen HLA-A11 alpha chain precursor - human

C:Species: Homo sapiens (man)
 C>Date: 31-Dec-1993 #sequence_revision 28-Apr-1995 #text_change 23-Jul-1999
 C:Accession: S03536; S03694; A47636; I60129
 R:Mayer, W.E.; Jonker, M.; Klein, D.; Ivanyi, P.; van Seventer, G.; Klein, J.
 EMBO J. 7, 2765-2774, 1988

A>Title: Nucleotide sequences of chimpanzee MHC class I alleles: evidence for trans-spec
 A:Reference number: S01171; MUID:89030641; PMID:2460344

A:Accession: S03536
 A:Molecule type: mRNA

A:Residues: 1-365 <MAY>

A:Cross-references: EMBL:X13111; NID:G32138; PIDN:CAA31503.1; PID:G32139
 A>Note: this allele is designated A*1101 (formerly All.1)

A:Accession: S03694
 A:Molecule type: mRNA

A:Residues: 1-42, 'K', 44-298 <MA2>

A:Cross-references: EMBL:X13112; NID:G32142; PIDN:CAA31504.1; PID:G32143
 A>Note: this allele is designated A*1102 (formerly All.1, All.2)

R:Cowan, E.P.; Jelachich, M.L.; Biddison, W.E.; Colligan, J.E.
 Immunogenetics 25, 241-250, 1987

A>Title: DNA sequence of HLA-A11: remarkable homology with HLA-A3 allows identification
 A:Reference number: A47636; MUID:87192928; PMID:2437024

A:Accession: A47636
 A:Molecule type: DNA

A:Residues: 26-365 <COW>

A:Cross-references: GB:M16007; GB:M16008; GB:M16009; GB:M16010; NID:G184130; PIDN:AAA654
 A>Note: the authors translated the codon GAC for residue 89 as Ala, CCG for residue 104

A>Note: this allele is designated A*1101 (formerly All.1)
 R:Lin, L.; Tokunaga, K.; Ishikawa, Y.; Bannai, M.; Kashiwase, K.; Kuwata, S.; Akaza, T.;
 Tissue Antigens 43, 78-82, 1994
 A>Title: Sequence analysis of serological HLA-A11 split antigens, All.1 and All.2.
 A:Reference number: I60129; MUID:94287401; PMID:8016845

A:Accession: I60129

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: mRNA

A:Residues: 1-365 <RES>

A:Cross-references: GB:D16841; NID:G540516; PIDN:BA04117.1; PID:G487909
 A>Note: this allele is designated A*1101 (formerly All.1)

C:Genetics:

A:Gene: GDB:HLA-A

A:Cross-references: GDB:119310; OMIM:142800

A:Map position: 6p21.3-6p21.3

C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
 C:Keywords: transmembrane protein

F;1-24/Domain: signal sequence #status predicted <SIG>

F;25-365/Product: class I histocompatibility antigen alpha chain #status predicted <EXT>

F;220-285/Domain: extracellular #status predicted <IMM>

F;220-285/Domain: immunoglobulin homology <IMM>

F;299-337/Domain: transmembrane #status predicted <TM>

F;338-365/Domain: intracellular #status predicted <INT>

Query Match 33.8%; Score 514; DB 2; Length 365;
 Best Local Similarity 39.4%; Pred. No. 3e-35;
 Matches 109; Conservative 47; Mismatches 113; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDE--SRVPERTPWVSSRISSQW 62

DB 26 SHSMRYFYTSVRGCRGPRFIAVGYDDTQVRFDSDAASQRMPEAPWIEQE-GPEYW 84

QY 63 LQLSLSLQKGDHMFVDFWTIMENHNHKE-SHTLQVILGCEMQEDNS-TEGWYKYGYDG 120

DB 85 DQETRNVAQSDTRVDLGLTRGYNQSDGSHTIQIMYGCDDVPGDGRFLGRQDAYDG 144

QY 121 QHLEFCPDTLDWRAAPRAWPKLEWERHKIRARONRAYLERDCPAQLQQLLELGRVL 180

DB 145 KDYALNEDLRSWTAADMAAQITKRWAAH-AAEQRAYLEGRVLEWLRYLENGKETL 203

QY 181 DQVPPPLVKVTHH-VTSSVTLRCALNYYPNITMKWLKDKQPMDAKEPEPKDVLNPD 239

DB 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTVELVETRPAGD 262

QY 240 GTYQGWITLAVPPGEORYTCQVHPGLDPLIVWE 276

DB 263 GTFQKAAVVVPSGGEORYTCHVQHEGLPKPLTLRWE 299

RESULT 10

HLA-A30.3 precursor - human

C:Species: Homo sapiens (man)

C>Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 21-Jan-2000

C:Accession: I56039

R:Kato, K.; Trapani, J.A.; Alloupena, J.; Dupont, B.; Yang, S.Y.

J. Immunol. 143, 3371-3378, 1989

A>Title: Molecular analysis of the serologically defined HLA-Aw19 antigens. A genetical.

A:Reference number: I56039; MUID:90038496; PMID:2478623

A:Accession: I56039

A:Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: DNA

A:Residues: 1-365 <RES>

A:Cross-references: GB:M30576; NID:G187646; PIDN:AAA59612.1; PID:G386878

C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.8%; Score 514; DB 2; Length 365;
 Best Local Similarity 39.4%; Pred. No. 3e-35;
 Matches 109; Conservative 48; Mismatches 112; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDE--SRVPERTPWVSSRISSQW 62

DB 26 SHSMRYFYTSVRGCRGPRFIAVGYDDTQVRFDSDAASQRMPEAPWIEQE-GPEYW 84

QY 63 LQLSLSLQKGDHMFVDFWTIMENHNHKE-SHTLQVILGCEMQEDNS-TEGWYKYGYDG 120

DB 85 DQETRNVAQSDTRVDLGLTRGYNQSDGSHTIQIMYGCDDVPGDGRFLGRQDAYDG 144

QY 121 QHLEFCPDTLDWRAAPRAWPKLEWERHKIRARONRAYLERDCPAQLQQLLELGRVL 180

DB 145 KDYALNEDLRSWTAADMAAQITKRWAAH-AAEQRAYLEGRVLEWLRYLENGKETL 203

QY 181 DQVPPPLVKVTHH-VTSSVTLRCALNYYPNITMKWLKDKQPMDAKEPEPKDVLNPD 239

DB 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTVELVETRPAGD 262

QY 240 GTYQGWITLAVPPGEORYTCQVHPGLDPLIVWE 276

DB 263 GTFQKAAVVVPSGGEORYTCHVQHEGLPKPLTLRWE 299


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Db 26 SHSMRYFTSVSRGSGEPRIAGVYDDTQFVRFSDAASQRMPEPAPWIEQE-RPEYW 84
Qy 63 LQLSQSLKGDHMTFVDFWTIMENHNHSE-SHTLQVILGCEMQEDNS--TEGYWKYGYDG 120
Db 85 DQETRNKAQSQTRVDLGLTGLRGYNQSEAGSHTIQIMYGCVDGSDGRFLRGYEQHAYDG 144
Qy 121 QDHLFECPTLDWRAAPRAWPPTKLEWRHKIRARQNRAYLDRDCPAQLQQLLELGRGVL 180
Db 145 KDYIALNEDLRSWTAAADMAAQITKRWAAAR-WAEQLRAYLEGTCVEWLRVLENGKETL 203
Qy 181 DQGVPLPVKVTHH-VTSSVTLRCALNYYPQNTIMKWLKDKQPMDAKEFEPEKDVLPNGD 239
Db 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLVELVETRPAGD 262
Qy 240 GTYQGWITLAVPPEGEORYTCQVEHPGLDQPLVINE 276
Db 263 GTFQKAAVVPVSGEORYTCHVQHEGLPKPLTLRWE 299

RESULT 11
HLHUA3
MHC class I histocompatibility antigen HLA-A3 alpha chain precursor - human
C:Species: Homo sapiens (man)
C>Date: 17-Mar-1987 #sequence_revision 17-Mar-1987 #text_change 02-Sep-1997
C:Accession: A02192
R:Strachan, T.; Soderover, R.; Damotte, M.; Jordan, B.R.
EMBO J. 3, 887-894, 1984
A:Title: Complete nucleotide sequence of a functional class I HLA gene, HLA-A3: implicat
A:Reference number: A02192; MUID:84207948; PMID:6609814
A:Accession: A02192
A:Molecule type: DNA
A:Residues: 1-370 <STR>
C:Genetics:
A:Gene: GDB:HLA-A
A:Cross-references: GDB:119310; OMIM:142800
A:Map position: 6p21.3-6p21.3
A:Introns: 30/1; 120/1; 212/1; 304/1; 343/1; 354/1; 370/1
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
C:Keywords: duplication; glycoprotein; heterodimer; transmembrane protein; transplanta
F:1-29/Domain: signal sequence #status predicted <SIG>
F:30-312/Domain: class I histocompatibility antigen HLA-A3 alpha chain #status predicted
F:30-312/Domain: extracellular #status predicted <EXT>
F:30-119/Domain: alpha-1 <EX1>
F:120-211/Domain: alpha-2 <EX2>
F:225-290/Domain: immunoglobulin homology <IMM>
F:313-337/Domain: transmembrane #status predicted <TM>
F:338-370/Domain: intracellular #status predicted <INT>
F:115/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:232-288/Disulfide bonds: #status predicted

Query Match 33.7%; Score 512; DB 1; Length 370;
Best Local Similarity 39.6%; Pred. No. 4.5e-35;
Matches 110; Conservative 47; Mismatches 111; Indels 10; Gaps 8;

Qy 5 SHSLHYLFMGASQDGLSLFALGVYDDQLFVYDDE--SRVPRTPWSSRSIQMW 62
Db 31 SHSMRYFTSVSRGSGEPRIAGVYDDTQFVRFSDAASQRMPEPAPWIEQE-GPEYW 89

Qy 63 LQLSQSLKGDHMTFVDFWTIMENHNHSE-SHTLQVILGCEMQEDNS--TEGYWKYGYDG 120
Db 90 DQETRNKAQSQTRVDLGLTGLRGYNQSEAGSHTIQIMYGCVDGSDGRFLRGYEQHAYDG 149
Qy 121 QDHLFECPTLDWRAAPRAWPPTKLEWRHKIRARQNRAYLDRDCPAQLQQLLELGRGV 179
Db 150 KDYIALNEDLRSWTAAADMAAQITKRWAAAR-WAEQLRAYLEGTCVEWLRVLENGKET 207
Qy 180 LQGVPLPVKVTHH-VTSSVTLRCALNYYPQNTIMKWLKDKQPMDAKEFEPEKDVLPNG 238
Db 208 LQRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLVELVETRPAG 266
Qy 239 DGTGQWITLAVPPEGEORYTCQVEHPGLDQPLVINE 276
Db 267 DGTGQWAAVVPVSGEORYTCHVQHEGLPKPLTLRWE 304
```

RESULT 12

I38439

MHC class I histocompatibility antigen HLA-A*8001 precursor - human

C:Species: Homo sapiens (man)

C>Date: 07-Jun-1996 #sequence_revision 07-Jun-1996 #text_change 21-Jan-2000

C:Accession: I59638; I38439

R:Domena, J.D.; Hildebrand, W.H.; Bias, W.B.; Parham, P.

Tissue Antigens 42, 156-159, 1993

A:Title: A sixth family of HLA-A alleles defined by HLA-A*8001.

A:Reference number: I59638; MUID:94112691; PMID:8284791

A:Accession: I59638

A:Status: preliminary; translated from GB/EMBL/DBDJ

A:Molecule type: mRNA

A:Residues: 1-365 <DOM>

R:Balas, A.; Garcia-Sanchez, F.; Gomez-Reino, F.; Vicario, J.L.

Immunogenetics 39, 452, 1994

A:Title: Characterization of a new and highly distinguishable HLA-A allele in a Spanish

A:Reference number: I38439; MUID:94245293; PMID:8188325

A:Accession: I38439

A:Status: preliminary; translated from GB/EMBL/DBDJ

A:Molecule type: mRNA

A:Residues: 1-365 <BAL>

A:Cross-references: EMBL:U03754; NID:g4322407; PIDN:AAC04322.1; PID:g4322408

C:Genetics:

A:Gene: GDB:HLA-A

A:Cross-references: GDB:119310; OMIM:142800

A:Map position: 6p21.3-6p21.3

C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

F:220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.6%; Score 510; DB 2; Length 365;

Best Local Similarity 38.3%; Pred. No. 6.5e-35;

Matches 106; Conservative 53; Mismatches 110; Indels 8; Gaps 7;

Qy 5 SHSLHYLFMGASQDGLSLFALGVYDDQLFVYDDE--SRVPRTPWSSRSIQMW 62

Db 26 SHSMRYFTSVSRGSGEPRIAGVYDDTQFVRFSDAASQRMPEPAPWIEQE-RPEYW 84

Qy 63 LQLSQSLKGDHMTFVDFWTIMENHNHSE-SHTLQVILGCEMQEDNS--TEGYWKYGYDG 120

Db 85 DEETRNKAKHSQTNRAHLGLTGLRGYNQSDGSHTIQIMYGCVDGSDGRFLRGYQDAYDG 144

Qy 121 QDHLFECPTLDWRAAPRAWPPTKLEWRHKIRARQNRAYLDRDCPAQLQQLLELGRGVL 180

Db 145 KDYIALNEDLRSWTAAADMAAQITKRWAAAR-WAEQLRAYLEGTCVEWLRVLENGKETL 203

Qy 181 DQGVPLPVKVTHH-VTSSVTLRCALNYYPQNTIMKWLKDKQPMDAKEFEPEKDVLPNGD 239

Db 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLVELVETRPAGD 262

Qy 240 GTYQGWITLAVPPEGEORYTCQVEHPGLDQPLVINE 276

Db 263 GTFQKAAVVPVSGEORYTCHVQHEGLPKPLTLRWE 299

RESULT 13

I37542

MHC class I histocompatibility antigen HLA-A2 alpha chain (allele A*0216) precursor - h

C:Species: Homo sapiens (man)

C>Date: 04-Oct-1996 #sequence_revision 04-Oct-1996 #text_change 21-Jan-2000

C:Accession: I37542; S49582

R:Barouch, D.; Krausa, P.; Bodmer, J.; Browning, M.J.; McMichael, A.J.

Immunogenetics 41, 388, 1995

A:Title: Identification of a novel HLA-A2 subtype, HLA-A*0216.

A:Reference number: I37542; MUID:95278976; PMID:7759139

A:Accession: I37542

A:Status: preliminary; translated from GB/EMBL/DBDJ

A:Molecule type: mRNA

A:Residues: 1-365 <RES>

A:Cross-references: EMBL:Z46633; NID:g575248; PIDN:CAA86602.1; PID:g575249

A;Note: submitted to the EMBL Data Library, November 1994

C;Genetics:

A;Gene: hla-A

C;Species: Homo sapiens (man)

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.5%; Score 509; DB 2; Length 365;
Best Local Similarity 39.4%; Pred. No. 7.9e-35;
Matches 109; Conservative 46; Mismatches 114; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDGLSLFEALGYVDDQLFVYDDE--SRVPRTPWSSRISSQMW 62
DB 26 SHSMRYFTTSVRSGRGEPRFIAVGYDDTQVRFDSDAASQRMPEPRAPIEQE-GPEYW 84
QY 63 LQLSLSLKGDHMFVTDFWTIMENHNHNSKE-SHTLQVILGCEMQED-NSTEGYWKYGYDG 120
DB 85 DGETRKVKAKHSQTHRVDLGLTRGYNQSEAGSHTVQRMVCGDVGSDWRFLRGYHQYAYDG 144
QY 121 QDHLFCFCDPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 KDVIYALKEDLRSWTAADMAAQTTHKWEAAHV-AEQLRAYLEGCEVWLRRLYLENGKETL 203
QY 181 DQOVPLVVKVTHH-VTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEFEKDVLPNGD 239
DB 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPYPAEITLTWQRDGED-QTQDTLVELTRPAGD 262
QY 240 GTYQGWITTLAVPPEGEORYTCQVEHPGLDQPLIVWE 276
DB 263 GTFOKWAADVVPVSGGEORYTCHVQHEGLPKPLTLRWE 299

RESULT 14

I38442

gene HLA-A-0205 protein - human

C;Species: Homo sapiens (man)

C;Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 21-Jan-2000

C;Accession: I38442

J;Immunol. 139, 936-941, 1987

A;Title: Multiple genetic mechanisms have contributed to the generation of the HLA-A2/A23

A;Reference number: I38441; MUID: 87252273; PMID: 3496393

A;Accession: I38442

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-365 <RES>

A;Cross-references: EMBL:U03862; NID:9432436; PIDN:AAA03603.1; PID:9432437

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.5%; Score 509; DB 2; Length 365;
Best Local Similarity 39.7%; Pred. No. 7.9e-35;
Matches 110; Conservative 44; Mismatches 115; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDGLSLFEALGYVDDQLFVYDDE--SRVPRTPWSSRISSQMW 62
DB 26 SHSMRYFTTSVRSGRGEPRFIAVGYDDTQVRFDSDAASQRMPEPRAPIEQE-GPEYW 84
QY 63 LQLSLSLKGDHMFVTDFWTIMENHNHNSKE-SHTLQVILGCEMQED-NSTEGYWKYGYDG 120
DB 85 DGETRKVKAKHSQTHRVDLGLTRGYNQSEAGSHTVQRMVCGDVGSDWRFLRGYHQYAYDG 144
QY 121 QDHLFCFCDPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 KDVIYALKEDLRSWTAADMAAQTTHKWEAAHV-AEQLRAYLEGCEVWLRRLYLENGKETL 203
QY 181 DQOVPLVVKVTHH-VTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEFEKDVLPNGD 239
DB 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPYPAEITLTWQRDGED-QTQDTLVELTRPAGD 262
QY 240 GTYQGWITTLAVPPEGEORYTCQVEHPGLDQPLIVWE 276
DB 263 GTFOKWAADVVPVSGGEORYTCHVQHEGLPKPLTLRWE 299

RESULT 15

I61902

MHC class I histocompatibility antigen HLA-A alpha chain precursor - human (isolate A*0212)

C;Species: Homo sapiens (man)

A;Variety: isolate A*0212

C;Date: 06-Sep-1996 #sequence_revision 06-Sep-1996 #text_change 23-Jul-1999

C;Accession: I61902

R;Belich, M.P.; Madrigal, J.A.; Hildebrand, W.H.; Zemmour, J.; Williams, R.C.; Luz, R.;

Nature 357, 326-329, 1992

A;Title: Unusual HLA-B alleles in two tribes of Brazilian Indians.

A;Reference number: I37120; MUID:92269955; PMID:1317015

A;Accession: I61902

A;Status: translated from GB/EMBL/DBJ

A;Molecule type: mRNA

A;Residues: 1-365 <RES>

A;Cross-references: GB:M84378; NID:9187625; PIDN:AAA59604.1; PID:9187626

A;Experimental source: cell line KRC 033; isolate A*0212

C;Genetics:

A;Gene: GDB:HLA-A

A;Cross-references: GDB:119310; OMIM:142800

A;Map position: 6p21.3-6p21.3

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

C;Keywords: transmembrane protein

F;1-24/Domain: signal sequence #status predicted <SIG>

F;25-365/Product: MHC class I histocompatibility antigen HLA-A alpha chain #status pred.

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.5%; Score 509; DB 2; Length 365;
Best Local Similarity 39.4%; Pred. No. 7.9e-35;
Matches 109; Conservative 45; Mismatches 115; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDGLSLFEALGYVDDQLFVYDDE--SRVPRTPWSSRISSQMW 62
DB 26 SHSMRYFTTSVRSGRGEPRFIAVGYDDTQVRFDSDAASQRMPEPRAPIEQE-GPEYW 84
QY 63 LQLSLSLKGDHMFVTDFWTIMENHNHNSKE-SHTLQVILGCEMQED-NSTEGYWKYGYDG 120
DB 85 DGETRKVKAKHSQTHRVDLGLTRGYNQSEAGSHTVQRMVCGDVGSDWRFLRGYHQYAYDG 144
QY 121 QDHLFCFCDPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 KDVIYALKEDLRSWTAADMAAQTTHKWEAAHV-AEQLRAYLEGCEVWLRRLYLENGKETL 203
QY 181 DQOVPLVVKVTHH-VTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEFEKDVLPNGD 239
DB 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPYPAEITLTWQRDGED-QTQDTLVELTRPAGD 262
QY 240 GTYQGWITTLAVPPEGEORYTCQVEHPGLDQPLIVWE 276
DB 263 GTFOKWAADVVPVSGGEORYTCHVQHEGLPKPLTLRWE 299

Search completed: August 5, 2003, 13:10:36

Job time : 16 secs

[7] SEQUENCE FROM N.A. (ISOFORMS 1; 7; 8; 9 AND 10).
 RP MEDLINE=20448010; PubMed=11001625;
 RX Thénie A., Orhant M., Gicquel I., Fergelot P., Le Gall J.-Y.,
 RA David V., Mosser J.;
 RT "The HFE gene undergoes alternate splicing processes.";
 RL Blood Cells Mol. Dis. 26:155-162(2000).
 RN [8]
 RP FUNCTION.
 RP MEDLINE=98132614; PubMed=9465039;
 RX Feder J.N., Penny D.M., Irfink A., Lee V.K., Lebron J.A., Watson N.,
 RA Teuchihaishi Z., Sigal E., Bjorkman P.J., Schatzman R.C.;
 RT "The hemochromatosis gene product complexes with the transferrin
 receptor and lowers its affinity for ligand binding.";
 RL Proc. Natl. Acad. Sci. U.S.A. 95:1472-1477(1998).
 RN [9]
 RP X-RAY CRYSTALLOGRAPHY (2.6 ANGSTROMS).
 RP MEDLINE=98206473; PubMed=9546397;
 RX Lebron J.A., Bennett M.J., Vaughn D.E., Chirino A.J., Snow P.M.,
 RA Mantler G.A., Feder J.N., Bjorkman P.J.;
 RT "Crystal structure of the hemochromatosis protein HFE and
 characterization of its interaction with transferrin receptor.";
 RL Cell 93:111-123(1998).
 RN [10]
 RP VARIANTS HH ASP-63 AND TYR-282.
 RX MEDLINE=97260408; PubMed=9106528;
 RA Carella M., D'Ambrosio L., Totaro A., Grifa A., Valentino M.A.,
 RA Piperno A., Girelli D., Roetto A., Franco B., Gasparini P.,
 RA Camaschella C.;
 RT "Mutation analysis of the HLA-H gene in Italian hemochromatosis
 patients.";
 RL Am. J. Hum. Genet. 60:828-832(1997).
 RN [11]
 RP VARIANT HH/PCT TYR-282.
 RX MEDLINE=97176837; PubMed=9024376;
 RA Roberts A.G., Whitley S.D., Morgan R.R., Worwood M., Elder G.H.;
 RT "Increased frequency of the hemochromatosis Cys282Tyr mutation in
 RT sporadic porphyria cutanea tarda.";
 RL Lancet 349:321-323(1997).
 RN [12]
 RP VARIANTS HH/PCT ASP-63.
 RX MEDLINE=98085904; PubMed=9425935;
 RA Sampietro M., Piperno A., Lupica L., Arosio C., Vergani A.,
 RA Corbetta N., Malosio I., Mattioli M., Fracanzani A.L.,
 RA Cappellini M.D., Fiorelli G., Fargion S.;
 RT "High prevalence of the His63Asp HFE mutation in Italian patients with
 RT porphyria cutanea tarda.";
 RL Hepatology 27:181-184(1998).
 RN [13]
 RP VARIANTS HH/PCT ASP-63 AND TYR-282.
 RX MEDLINE=98281650; PubMed=9620340;
 RA Bonkovsky H.L., Poh-Fitzpatrick M., Pinestone N., Obando J.,
 RA Di Biasele A., Tattire C., Tortorelli K., LeClair P., Mercurio M.G.,
 RA Lambercht R.W.;
 RT "Porphyria cutanea tarda, hepatitis C, and HFE gene mutations in North
 RT America.";
 RL Hepatology 27:1661-1669(1998).
 RN [14]
 RP VARIANTS HH ASP-63; CYS-65 AND TYR-282.
 RX MEDLINE=99211934; PubMed=10194428;
 RA Mura C., Raqueno O., Ferec C.;
 RT "HFE mutations analysis in 711 hemochromatosis probands: evidence for
 RT S65C implication in mild form of hemochromatosis.";
 RL Blood 93:2502-2505(1999).
 RN [15]
 RP VARIANTS HH CYS-65; ARG-93 AND THR-105.
 RX MEDLINE=20042794; PubMed=10575540;
 RA Barton J.C., Sawada-Hirai R., Rothenberg B.E., Acton R.T.;
 RT "Two novel missense mutations of the HFE gene (I105T and G93R) and
 RT identification of the S65C mutation in Alabama hemochromatosis
 RT probands.";
 RL Blood Cells Mol. Dis. 25:147-155(1999).
 RN [16]

RP VARIANTS VP ASP-63 AND HIS-127, VARIANT HH MET-330, AND VARIANTS
 RP MET-53 AND MET-59.
 RX MEDLINE=99330560; PubMed=10401000;
 RA de Villiers J.N.P., Hillermann R., Loubser L., Kotze M.J.;
 RT "Spectrum of mutations in the HFE gene implicated in haemochromatosis
 RT and porphyria.";
 RL Hum. Mol. Genet. 8:1517-1522(1999).
 RN [17]
 RP VARIANTS HH ASP-63 AND TYR-282.
 RX MEDLINE=99140360; PubMed=10094552;
 RA Merryweather-Clarke A.T., Simonsen H., Shearman J.D., Poynton J.J.,
 RA Norgaard-Pedersen B., Robson K.J.H.;
 RT "A retrospective anonymous pilot study in screening newborns for HFE
 RT mutations in Scandinavian populations.";
 RL Hum. Mutat. 13:154-159(1999).
 RN [18]
 RP VARIANT HH CYS-65.
 RX Fagan E., Payne S.J.;
 RA "A novel missense mutation S65C in the HFE gene with a possible role
 RT in hereditary haemochromatosis.";
 RL Hum. Mutat. 13:507-508(1999).
 RN [19]
 RP VARIANT LYS-277.
 RX MEDLINE=20081073; PubMed=10612845;
 RA Bradbury R., Fagan E., Payne S.J.;
 RT "Two novel polymorphisms (E277K and V212V) in the haemochromatosis
 RT gene HFE.";
 RL Hum. Mutat. 15:120-120(2000).
 CC -1- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 CC affinity for iron-loaded transferrin.
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=10;
 CC Comment=Additional isoforms seem to exist;
 CC Name=1;
 CC IsoId=Q30201-1; Sequence=Displayed;
 CC Name=2; Synonyms=delE2;
 CC IsoId=Q30201-2; Sequence=VSP_003218;
 CC Name=3; Synonyms=del14E4;
 CC IsoId=Q30201-3; Sequence=VSP_003225;
 CC Name=4; Synonyms=delE214E4;
 CC IsoId=Q30201-4; Sequence=VSP_003218, VSP_003225;
 CC Name=5;
 CC IsoId=Q30201-5; Sequence=VSP_003219;
 CC Name=6;
 CC IsoId=Q30201-6; Sequence=VSP_003220;
 CC Name=7; Synonyms=delE3;
 CC IsoId=Q30201-7; Sequence=VSP_003221;
 CC Name=8; Synonyms=1043-2283del,intron6ins;
 CC IsoId=Q30201-8; Sequence=VSP_003226, VSP_003227;
 CC Name=9; Synonyms=delE3-7;
 CC IsoId=Q30201-9; Sequence=VSP_003223, VSP_003224;
 CC Name=10; Synonyms=582-878del;
 CC IsoId=Q30201-10; Sequence=VSP_003222;
 CC -1- TISSUE SPECIFICITY: IN ALL TISSUES TESTED EXCEPT BRAIN.
 CC -1- DISEASE: DEFECTS IN HFE ARE A CAUSE OF HEREDITARY HEMOCHROMATOSIS
 CC (HH). HH IS AN AUTOSOMAL RECESSIVE INBORN DISORDER OF IRON
 CC METABOLISM. FREQUENT AMONG CAUCASIANS. HH IS CHARACTERIZED BY
 CC ABNORMAL INTESTINAL IRON ABSORPTION AND PROGRESSIVE INCREASE OF
 CC TOTAL BODY IRON, WHICH RESULTS IN MIDLIFE IN CLINICAL
 CC COMPLICATIONS INCLUDING CIRRHOSIS, CARDIOPATHY, DIABETES,
 CC ENDOCRINE DYSFUNCTIONS, ARTHROPATHY, AND SUSCEPTIBILITY TO LIVER
 CC CANCER. SINCE THE DISEASE COMPLICATIONS CAN BE EFFECTIVELY
 CC PREVENTED BY REGULAR PHLEBOTOMIES, EARLY DIAGNOSIS IS MOST
 CC IMPORTANT TO PROVIDE A NORMAL LIFE EXPECTANCY TO THE AFFECTED
 CC SUBJECTS.
 CC -1- DISEASE: DEFECTS IN HFE ARE A CAUSE OF PORPHYRIA CUTANEA TARDA
 CC (PCT), A DISORDER CHARACTERIZED BY LIGHT-SENSITIVE DERMATITIS AND
 CC PRESENCE OF LARGE AMOUNTS OF UROPORPHYRIN IN URINE. IRON OVERLOAD
 CC IS OFTEN PRESENT IN ASSOCIATION WITH VARYING DEGREES OF LIVER
 CC DAMAGE. PCT IS THE MOST COMMON FORM OF PORPHYRIA WORLDWIDE. IT
 CC OCCURS IN TWO FORMS: THE SPORADIC TYPE (PCT TYPE I) AND THE
 CC FAMILIAL TYPE (PCT TYPE II), WHICH IS INHERITED IN AN AUTOSOMAL

Query Match 99.5%; Score 1513; DB 1; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.3e-119;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFFVYDDSRVPRTPWSSRSISQ 60
DB 23 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFFVYDDSRVPRTPWSSRSISQ 82

QY 61 MWLQSLKSGWDMFTVDFWMTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLKSGWDMFTVDFWMTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142

QY 121 ODHLEFCDDTLDRAAEPRAWPTKLEWERHKKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 ODHLEFCDDTLDRAAEPRAWPTKLEWERHKKIRARQNRAYLERDCPAQLQQLLELGRGVL 202

QY 181 DQVPPVLKVTHTVTSVTLRCALNYPQNTMKWLKDKQPMDAKEFEPEKDVLPNGD 240
DB 203 DQVPPVLKVTHTVTSVTLRCALNYPQNTMKWLKDKQPMDAKEFEPEKDVLPNGD 262

QY 241 TYQGWITLAVPPGEGORYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPPGEGORYTCQVEHPGLDQPLIVWE 298

RESULT 2
HFE DICSU STANDARD; PRT; 348 AA.

AC Q9GL42;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hereditary hemochromatosis protein precursor.
GN HFE.

OS Dicerorhinus sumatrensis (Sumatran rhinoceros).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Dicerorhinus.
OX NCBI_TaxID=89632;
RN [1]
RP SEQUENCE FROM N.A.
RA West C.J., Worley M., Beutler E.;
RT "Rhinoceros HFE polymorphisms."
RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
CC affinity for iron-loaded transferrin.
CC -!- SUBCELLULAR LOCATION: Type I membrane protein.
CC -!- SIMILARITY: TO MHC CLASS I ANTIGENS.

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CC or send an email to license@isb-sib.ch).

CC EMBL; AY007543; AAG23703.1; -.
CC HSSP; Q30201; 1A6Z.
CC InterPro; IPR007110; Ig-like.
CC InterPro; IPR003597; Ig-cl.
CC InterPro; IPR003006; Ig_MHC.
CC InterPro; IPR001039; MHC_I.
CC Pfam; PF00047; Ig_1.
CC Pfam; PF00129; MHC_I; 1.
CC ProDom; PD000050; MHC_I; 1.
CC SMART; SM00407; IGC1; 1.
CC PROSITE; PS50835; IG_LIKE; 1.
CC PROSITE; PS00290; IG_MHC; 1.
CC MHC_I; Transmembrane; Glycoprotein; Signal.
KW MHC_I; 22 BY SIMILARITY.
FT SIGNAL 1 22
FT CHAIN 23 348 HEREDITARY HEMOCHROMATOSIS PROTEIN.
FT DOMAIN 23 114 EXTRACELLULAR ALPHA-1.

FT DOMAIN 115 205 EXTRACELLULAR ALPHA-2.
FT DOMAIN 206 297 EXTRACELLULAR ALPHA-3.
FT TRANSMEM 307 330 CONNECTING PEPTIDE.
FT TRANSMEM 331 348 POTENTIAL.
FT DISULFID 124 187 CYTOPLASMIC TAIL.
FT DISULFID 225 282 BY SIMILARITY.
FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 130 130 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 234 234 N-LINKED (GLCNAC. . .) (POTENTIAL).
SQ SEQUENCE 348 AA; 39740 MW; 518BFD357AB83B90 CRC64;

Query Match 81.4%; Score 1238; DB 1; Length 348;
Best Local Similarity 81.3%; Pred. No. 1.5e-96;
Matches 222; Conservative 20; Mismatches 31; Indels 0; Gaps 0;

QY 4 RSHSLHYLFMGASQDGLSLFEALGYVDQDLFFVYDDSRVPRTPWSSRSISQMWL 63
DB 26 RSHSLHYLFMGASQDGLSLFEALGYVDQDLFFVYDDSRVPRTPWSSRSISQMWL 85

QY 64 QLSQSLKSGWDMFTVDFWMTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDGQDH 123
DB 86 QLSQSLKSGWDMFTVDFWMTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDGQDH 145

QY 124 LEFCDDTLDRAAEPRAWPTKLEWERHKKIRARQNRAYLERDCPAQLQQLLELGRGVL 183
DB 146 LEFCDDTLDRAAEPRAWPTKLEWERHKKIRARQNRAYLERDCPAQLQQLLELGRGVL 205

QY 184 VPPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEFEPEKDVLPNGDGTQ 243
DB 206 VPPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEFEPEKDVLPNGDGTQ 265

QY 244 GWITLAVPPGEGORYTCQVEHPGLDQPLIVWE 276
DB 266 SWALAVPPGEGORYTCQVEHPGLDQPLIVWE 298

RESULT 3
HFE CERSI STANDARD; PRT; 348 AA.

AC Q9GKZ0;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hereditary hemochromatosis protein precursor.
GN HFE.

OS Ceratotherium simum (White rhinoceros) (Square-lipped rhinoceros).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Ceratotherium.
OX NCBI_TaxID=9807;
RN [1]
RP SEQUENCE FROM N.A.
RA West C.J., Worley M., Beutler E.;
RT "Rhinoceros HFE polymorphisms."
RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
CC affinity for iron-loaded transferrin.
CC -!- SUBCELLULAR LOCATION: Type I membrane protein.
CC -!- SIMILARITY: TO MHC CLASS I ANTIGENS.

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CC or send an email to license@isb-sib.ch).

CC EMBL; AY007541; AAG23701.1; -.
CC HSSP; Q30201; 1A6Z.
CC InterPro; IPR007110; Ig-like.
CC InterPro; IPR003597; Ig-cl.
CC InterPro; IPR003006; Ig_MHC.
CC InterPro; IPR001039; MHC_I.
CC Pfam; PF00047; Ig_1.
CC Pfam; PF00129; MHC_I; 1.
CC ProDom; PD000050; MHC_I; 1.
CC SMART; SM00407; IGC1; 1.
CC PROSITE; PS50835; IG_LIKE; 1.
CC PROSITE; PS00290; IG_MHC; 1.
CC MHC_I; Transmembrane; Glycoprotein; Signal.
KW MHC_I; 22 BY SIMILARITY.
FT SIGNAL 1 22
FT CHAIN 23 348 HEREDITARY HEMOCHROMATOSIS PROTEIN.
FT DOMAIN 23 114 EXTRACELLULAR ALPHA-1.


```

QY 5 SHSLHYLFMGASEODLGLSFEALGYDDQLFVYDDERRRPRTPWVSSRISQWLQ 64
DB 32 SHSLRYLFMGASRPDLGLPFEALGYDDQLFVSYNHSRAEPRAPIWLGQTSSQLWLQ 91
QY 65 LSQSLKGDHMTVDFTWIMENHNSK-----ESHTLQVILGCEMQEDNSTEGWKY 116
DB 92 LSQSLKGDHMTVDFTWIMENHNSK-----ESHTLQVILGCEMQEDNSTEGWKY 151
QY 117 GYGQDHLFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELG 176
DB 152 GYGQDHLFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELG 211
QY 177 RGVLQDQVPLVKVTHVTSVTLRCALNYYPQNTIMKWKDKQPMDAKEPEPKDVL 236
DB 212 RGVLQDQVPLVKVTHVTSVTLRCALNYYPQNTIMKWKDKQPMDAKEPEPKDVL 271
QY 237 NGDGTQGWTLAVPQGEORYTCQVHPGLDQPLVIWE 276
DB 272 NGDGTQGWTLAVPQGEORYTCQVHPGLDQPLVIWE 311

RESULT 7
HFE_MOUSE
ID HFE_MOUSE STANDARD; PRT; 359 AA.
AC P70387;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hereditary hemochromatosis protein homolog precursor.
GN HFE OR MR2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RN SEQUENCE FROM N.A.
RC STRAIN=129/SvJ;
RX MEDLINE=98060831; PubMed=9396865;
RA Riegert P., Gillilan S., Nanda I., Schmid M., Bahram S.;
RT "The mouse HFE gene.";
RL Immunogenetics 47:174-177 (1998).
RN [2]
RN SEQUENCE FROM N.A.
RC STRAIN=BALB/c; TISSUE=Lung;
RA Hashimoto K.;
RL Submitted (SEP-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RN SEQUENCE OF 37-211 FROM N.A.
RC STRAIN=BALB/c; TISSUE=Liver;
RX MEDLINE=97148566; PubMed=9020055;
RA Hashimoto K., Hirai M., Kurosawa Y.;
RT "Identification of a mouse homolog for the human hereditary
hemochromatosis candidate gene.";
RL Biochem. Biophys. Res. Commun. 230:35-39 (1997).
RN [4]
RN SEQUENCE OF 79-359 FROM N.A.
RC STRAIN=129;
RA Albright W., Drabant B., Doenecke D.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
CC -I- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
affinity for iron-loaded transferrin (By similarity).
CC -I- SUBCELLULAR LOCATION: Type I membrane protein.
CC -----
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CC -----
EMBL; AF007558; AAC03447.1; -.

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DR EMBL; U66849; AAB07525.1; -.
DR EMBL; Y12650; CAA73197.1; -.
DR EMBL; U80604; AAB51504.1; -.
DR PIR; JCS382; JCS382.
DR HSSP; Q30201; IA62.
DR MGD; MGI:109191; Hfe.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_c1.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00447; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR PRINTS; PR01638; MHCCLASS1.
DR ProDom; PD000050; MHC I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS00835; IG-LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 359
FT HEREDITARY HEMOCHROMATOSIS PROTEIN
FT HOMOLOG.
FT DOMAIN 25 126
FT DOMAIN 127 217
FT DOMAIN 218 309
FT DOMAIN 310 318
FT TRANSMEM 319 339
FT TRANSMEM 340 359
FT DISULFID 136 139
FT DISULFID 237 294
FT CARBOHYD 114 142
FT CARBOHYD 142 142
FT CARBOHYD 166 166
FT CARBOHYD 246 246
FT SEQUENCE 359 AA; 40548 MW; 4BDE6C27F9FF20B4 CRC64;
SQ
Query Match 75.0%; Score 1140; DB 1; Length 359;
Best Local Similarity 72.2%; Pred. No. 2.5e-88;
Matches 203; Conservative 30; Mismatches 40; Indels 8; Gaps 1;
QY 4 RSHSLHYLFMGASEODLGLSFEALGYDDQLFVYDDERRRPRTPWVSSRISQWL 63
DB 30 RSHSLRYLFMGASEPDLGLPFEALGYDDQLFVSYNHSRAEPRAPIWLGQTSSQLWL 89
QY 64 QLSQSLKGDHMTVDFTWIMENHNSK-----ESHTLQVILGCEMQEDNSTEGWKY 115
DB 90 HLSQSLKGDHMTVDFTWIMENHNSK-----ESHTLQVILGCEMQEDNSTEGWKY 149
QY 116 GYGQDHLFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLEL 175
DB 150 GYGQDHLFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLEL 209
QY 176 GYGQDHLFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLEL 235
DB 210 GYGQDHLFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLEL 269
QY 236 PNGDGTQGWTLAVPQGEORYTCQVHPGLDQPLVIWE 276
DB 270 PNGDGTQGWTLAVPQGEORYTCQVHPGLDQPLVIWE 310
RESULT 8
HFE_MOUSE
ID HFE_MOUSE STANDARD; PRT; 361 AA.
AC P01894;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE RLA class I histocompatibility antigen, alpha chain 11/11 precursor.
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxID=9986;
RN [1]

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QY 121 QDHLFCPTDLNRAAPRAAPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
Db 145 ADVIALNEDLRSWTAADTAQNTQKWEAAG-EAERHRAYLERECVEWLRYLEMGKETL 203
QY 181 DQVPPPLVKTHTVSS-VTTLCRALNYPQNTMKWLKDKQPMDAKEEPEKDVLPNGD 239
Db 204 QRADPPKAVHTHPASDREATLRCWALGFYPAISLTWQDGDGSD-QTQDTLVELTRPGGD 262
QY 240 GTYQGWITLAVPPGEQRYTCQVHPGLDQPLIVIE 276
Db 263 GTFQKAAVVPVSGEQRYTCRVQHEGLPEPLTLTWE 299

RESULT 10
1A01 PANTR
ID 1A01 PANTR STANDARD; PRT; 365 AA.
AC P16209;
DT 01-APR-1990 (Rel. 14, Created)
DT 01-APR-1990 (Rel. 14, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE CHLA class I histocompatibility antigen, A-2 alpha chain precursor.
DE Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90201944; PubMed=1690682;
RA Lawlor D.A., Warren E., Ward F.E., Parham P.;
RT "Comparison of class I MHC alleles in humans and apes.";
RL Immunol. Rev. 113:147-185(1990).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
MICROGLOBULIN).
CC
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CC
CC EMBL; M30678; AAA87970.1; --
DR PIR; I36961; I36961.
DR HSP; Q95352; IHK.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS50835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
DR MHC_I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 365 CHLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
A-2 ALPHA CHAIN.
FT FT 25 114 EXTRACELLULAR ALPHA-1.
FT FT 25 206 EXTRACELLULAR ALPHA-2.
FT FT 207 298 EXTRACELLULAR ALPHA-3.
FT FT 299 308 CONNECTING PEPTIDE.
FT FT 308 332
FT TRANSMEM 332 365
FT DOMAIN 333 365
FT DOMAIN 333 365
FT DISULFID 125 188
FT FT 227 283 BY SIMILARITY.
FT FT 110 110 N-LINKED (GLCNAC...) (BY SIMILARITY).
FT CARBOHYD 365 AA; 40848 MW; FC452768D038D3E CRC64;
SQ SEQUENCE
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Query Match 34.0%; Score 517; DB 1; Length 365;
Best Local Similarity 39.7%; Pred. No. 3.9e-36;
Matches 110; Conservative 45; Mismatches 114; Indels 8; Gaps 7;

QY 5 SLSHLVFMGASQDGLSLFEALGVVDQLFVFDDE--SRVERPTWVSSRISSQMW 62
Db 26 SLSMRVFFTSVSRGGEPRFIAVGVDVDTQVFRFSDAASQRMPEAPRWIEQ-GPEYW 84
QY 63 LQLSQLKGDHMFVTVDFTIMENHNHNSKE-SHTLQVILGCEMQEDNS-TEGYWYKTYDYG 120
Db 85 DEETRSKSAHSQTDTRVDLGLTGRGYNNQSGSHITQIMYGCDSGRFLGRVQDAYDG 144
QY 121 QDHLFCPTDLNRAAPRAAPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
Db 145 KDVIALLNEDLRSWTAADTAQNTQKWEAAG-EAERHRAYLERECVEWLRYLEMGKETL 203
QY 181 DQVPPPLVKTHTVSS-VTTLCRALNYPQNTMKWLKDKQPMDAKEEPEKDVLPNGD 239
Db 204 QRTDPEKTHMTHPTISDHEATLRCWALGFYPAISLTWQDGDGSD-QTQDTLVELTRPGD 262
QY 240 GTYQGWITLAVPPGEQRYTCQVHPGLDQPLIVIE 276
Db 263 GTFQKAAVVPVSGEQRYTCRVQHEGLPEPLTLTWE 299

RESULT 11
HAIB BOVIN STANDARD; PRT; 364 AA.
AC P13753;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE BOLA class I histocompatibility antigen, alpha chain BL3-7 precursor.
DE Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=88258075; PubMed=3133413;
RA Emnis P.D., Jackson A.P., Parham P.;
RT "Molecular cloning of bovine class I MHC cDNA.";
RL J. Immunol. 141:642-651(1988).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
MICROGLOBULIN).
CC
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CC
CC EMBL; M21043; AAA30641.1; --
DR HSP; P16391; LED3.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS50835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
DR MHC_I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 27
FT CHAIN 28 364 BOLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
A-2 ALPHA CHAIN.
```

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FT DOMAIN 28 117 ALPHA CHAIN BL3-7.
FT DOMAIN 118 209 EXTRACELLULAR ALPHA-1.
FT DOMAIN 210 301 EXTRACELLULAR ALPHA-2.
FT DOMAIN 302 310 EXTRACELLULAR ALPHA-3.
FT TRANSMEM 311 331 CONNECTING PEPTIDE.
FT DOMAIN 332 364 CYTOPLASMIC.
FT CARBOHYD 106 106 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 113 113 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT DISULFID 128 191 BY SIMILARITY.
FT DISULFID 230 286 BY SIMILARITY.
SQ SEQUENCE 364 AA; 41513 MW; 622056CF7DC67873 CRC64;

Query Match 33.8%; Score 516; DB 1; Length 364;
Best Local Similarity 38.9%; Pred. No. 4.8e-36;
Matches 109; Conservative 50; Mismatches 113; Indels 8; Gaps 7;

QY 2 LLRSLSLHLYFMGASEQDLGLSLFEALGYVDQDLFPYDDB--SRRVPRTPWSSRISS 59
DB 26 LAGSHSLRYPTGVSRGLGEPRIAGVYDDTQFVRFSDAPNPREEPRVPMWQE-GP 84
QY 60 QMWLQSLQSLKGDHMTVDFTWIMENHNHKSKE-SHTLQVILGCEMOEDNS-TEGYWKYG 117
DB 85 EYDNRTRIYKDTAQIFRVLNLTIRGYNQSETGSHNIQAMYGCDVCPDGLLGRFQFG 144
QY 118 YDQDHLFCFDPDLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQOLLELGR 177
DB 145 YDGRDYIALNEELRSWTAADTAQITKRKEAAG-AAETWRNLYEGECVWELRRYLENGK 203
QY 178 GVLDQVPPVPLVKTTH-VTSSTVTLRCALNYYPQNTMVKDKQPMDAKEPEPKDVL 236
DB 204 DTLRLADPPKAVHTHSISDREVTLCRWALGFYPEISLTWQREGED-QTDMLVETRP 262
QY 237 NGDGTQGMITLAVPGEEQRYTCQVEHPGLDPLIVIME 276
DB 263 SGGDTQKWAALVVPSEGEQRYTCRVQHEGLQEPDLRLWE 302

RESULT 12
ID 1A11 HUMAN STANDARD; PRT; 365 AA.
AC P13746;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE HLA class I histocompatibility antigen, A-11 alpha chain precursor.
GN HLA-A OR HLAA.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (A*1101/A*1102).
RX MEDLINE=89030641; PubMed=2460344;
RA Mayer W.B., Jonker M., Klein D., Ivanyi P., van Severen G.,
RA Klein J.;
RT "Nucleotide sequences of chimpanzee MHC class I alleles: evidence for
RT trans-species mode of evolution.";
RN EMBJ J. 7:2765-2774(1988).
RN [2]
RP SEQUENCE FROM N.A. (A*1101/A*1102).
RX MEDLINE=94287401; PubMed=8016845;
RA Lin L., Tokunaga K., Ishikawa Y., Bannai M., Kashiwase K.,
RA Kuwata S., Akaza T., Tadokoro K., Shibata Y., Juji T.;
RT "Sequence analysis of serological HLA-A11 split antigens, A11.1 and
RT A11.2.";
RN Tissue Antigens 43:78-82(1994).
RN [3]
RP SEQUENCE OF 26-365 FROM N.A. (A*1101).
RX MEDLINE=87192928; PubMed=2437024;
RA Cowan E.P., Jelachich M.L., Biddison W.E., Coligan J.E.;
RT "DNA sequence of HLA-A11: remarkable homology with HLA-A3 allows
RT identification of residues involved in epitopes recognized by
```

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RT antibodies and T cells.";
RL Immunogenetics 25:241-250(1987).
CC -I- FUNCTION: Involved in the presentation of foreign antigens to
CC the immune system.
CC -I- SUBUNIT: Dimer of alpha chain and a beta chain (beta-2-
CC microglobulin).
CC -I- POLYMORPHISM: THE FOLLOWING ALLELES OF A-11 ARE KNOWN: A*1101 (A-
CC 11E) AND A*1102 (A-11K). THE SEQUENCE SHOWN IS THAT OF A*1101.
CC
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CC
DR EMBL; X13111; CAA31503.1; -
DR EMBL; X13112; CAA31504.1; -
DR EMBL; D16841; BAA04117.1; -
DR EMBL; D16842; BAA04118.1; -
DR EMBL; M16010; AAA65449.1; -
DR EMBL; M16007; AAA65449.1; JOINED.
DR EMBL; M16008; AAA65449.1; JOINED.
DR EMBL; M16009; AAA65449.1; JOINED.
DR PIR; I83063; I83063.
DR PIR; S03536; A47636.
DR HSP; O19673; 1HSB.
DR MIM; 142800; -
DR GO; GO:0005887; C:integral to plasma membrane; NAS.
DR GO; GO:0030106; F:MHC class I receptor activity; NAS.
DR GO; GO:0006955; P:immune response; NAS.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; IG_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF0129; MHC I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IG1_1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal; Polymorphism.
FT SIGNAL 1 24
FT CHAIN 25 365 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT A-11 ALPHA CHAIN.
FT DOMAIN 25 114 EXTRACELLULAR ALPHA-1.
FT DOMAIN 115 206 EXTRACELLULAR ALPHA-2.
FT DOMAIN 207 298 EXTRACELLULAR ALPHA-3.
FT DOMAIN 299 308 CONNECTING PEPTIDE.
FT TRANSMEM 309 332
FT DOMAIN 333 365 CYTOPLASMIC TAIL.
FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT DISULFID 125 188 BY SIMILARITY.
FT DISULFID 227 283 BY SIMILARITY.
FT VARIANT 43 43 E -> K (IN ALLELE A*1102).
FT SEQUENCE 365 AA; 40937 MW; FE449CE2D4BF6CC5 CRC64;

Query Match 33.8%; Score 514; DB 1; Length 365;
Best Local Similarity 39.4%; Pred. No. 7e-36;
Matches 109; Conservative 47; Mismatches 113; Indels 8; Gaps 7;

QY 5 SHSLHLYFMGASEQDLGLSLFEALGYVDQDLFPYDDB--SRRVPRTPWSSRISSQW 62
DB 26 SHSMRYFTYSVSRPGEPRIAGVYDDTQFVRFSDAASQRMPEAPWISQE-GPEYV 84
QY 63 LQLSOSLKGWDHMTVDFTWIMENHNHKSKE-SHTLQVILGCEMOEDNS-TEGYWKYG 120
DB 85 DQETRNKVAQSQTDRVDLGLTIRGYNQSDGSHITQIMYGCDVCPDGLRGRQADVG 144
QY 121 QDHLFCFDPDLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQOLLELGR 180
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Db 145 KDVIALLNEDLSRTAADMAAQITKRWEAAH-AAEQRAYLEGRCVWELRRYLENGKETL 203
QY 181 DQOVPPPLVKVTH-VTSSVTTLCRALNYPQNTMKWLKDQPMADAKFEPKDVLPNGD 239
Db 204 QRTDPPKTHMTHPISDHEATLRCWALGFYFABITLTWQDGED-QTQDTLTVETRPAGD 262
QY 240 GTYQGMITLAVPGEQRQYTCQVEHGLDQPLIVWE 276
Db 263 GTFOKAAVVVPSGEGQRQYTCVQHEGLPKPLTLRWE 299

RESULT 13
ID 1A03 HUMAN STANDARD; PRT; 370 AA.
AC P04439;
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE HLA class I histocompatibility antigen, A-3 alpha chain precursor.
GN HLA-A OR HLA-A.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN 1;
RP SEQUENCE FROM N.A. (A*0301).
RX MEDLINE=84207948; PubMed=6609814;
RA Strachan T., Sodoyer R., Damotte M., Jordan B.R.;
RT "Complete nucleotide sequence of a functional class I HLA gene,
RT HLA-A3: implications for the evolution of HLA genes.";
RL EMBO J. 3:887-894 (1984).
RN 2;
RP SEQUENCE FROM N.A. (A*0302).
RX MEDLINE=85290871; PubMed=2993417;
RA Cowan E.P., Jordan B.E., Coligan J.E.;
RT "Molecular cloning and DNA sequence analysis of genes encoding
RT cytotoxic T lymphocyte-defined HLA-A3 subtypes: the E1 subtype.";
RL J. Immunol. 135:2835-2841 (1985).
CC -1- FUNCTION: Involved in the presentation of foreign antigens to
CC the immune system.
CC -1- SUBUNIT: Dimer of alpha chain and a beta chain (beta-2-
CC microglobulin).
CC -1- POLYMORPHISM: THE FOLLOWING ALLELES OF A-3 ARE KNOWN: A*0301 (A-
CC 3.1) AND A*0302. THE SEQUENCE SHOWN IS THAT OF A*0301.
CC -----
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CC -----
CC ENBL; X00492; CAA25162.1; ALT_TERM.
DR PIR; A02192; HLHUA3.
DR HSP; O19673; IHSB.
DR MIM; 142800;
DR GO; GO:0005887; C: integral to plasma membrane; NAS.
DR GO; GO:0030106; F: MHC class I receptor activity; NAS.
DR GO; GO:0006955; P: immune response; NAS.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_1.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PSS0835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; transmembrane; Glycoprotein; signal; Polymorphism.

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FT SIGNAL 1 29
FT CHAIN 30 370
FT FT HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT A-3 ALPHA CHAIN
FT DOMAIN 30 119 EXTRACELLULAR ALPHA-1.
FT DOMAIN 120 211 EXTRACELLULAR ALPHA-2.
FT DOMAIN 212 303 EXTRACELLULAR ALPHA-3.
FT DOMAIN 304 313 CONNECTING PEPTIDE.
FT TRANSMEM 314 337
FT DOMAIN 338 370 CYTOPLASMIC TAIL.
FT CARBOHYD 115 115 N-LINKED (GLUCNA. .) (BY SIMILARITY).
FT DISULFID 130 193 BY SIMILARITY.
FT DISULFID 232 288 E -> V (IN ALLELE A*0302).
FT VARIANT 181 181 /FTID=VAR 004351.
FT VARIANT 185 185 L -> Q (IN ALLELE A*0302).
FT /FTID=VAR 004352.
SQ SEQUENCE 370 AA; 41368 MW; ABLFA77460318A2 CRC64;
Query Match 33.7%; Score 512; DB 1; Length 370;
Best Local Similarity 39.6%; Pred. No. 1.1e-35;
Matches 110; Conservative 47; Mismatches 111; Indels 10; Gaps 8;
QY 5 SHSLHYLFMGAGEQDGLSLFALGVDDQLFVVDDE--SRREPTFWSSRISQMW 62
Db 31 SHSMRYFTTSVRPGRGEPRTIAGVYDVTQFVRFSDAASQRMPEAPWIEQ-GPEYW 89
QY 63 LQLSQSLKGDHMTFTVFWTMINHNHNSKE-SHTLQVILGCEMQEDNS-TEGYWKYGYDG 120
Db 90 DQETRNVAQSQSDTRVDLGLTGYNQSAGSHITQIMYGCDVGSDFRFLGRTRQDAYDG 149
QY 121 QHLSFCPTDLWRAAEPAWPTKLEWE--RHKIRARQNAYLERDCPAQLQQLLELGRGV 179
Db 150 KDVIALLNEDLSRTAADMAAQITKRWEAAH-AAEQRAYLDGTCVWELRRYLENGKET 207
QY 180 LDOQVPPPLVKVTH-VTSSVTTLCRALNYPQNTMKWLKDQPMADAKFEPKDVLPNG 238
Db 208 LQRTDPPKTHMTHPISDHEATLRCWALGFYFABITLTWQDGED-QTQDTLTVETRPAG 266
QY 239 DCTYQGMITLAVPGEQRQYTCQVEHGLDQPLIVWE 276
Db 267 DGTFOKAAVVVPSGEGQRQYTCVQHEGLPKPLTLRWE 304

RESULT 14
ID 1A80 HUMAN STANDARD; PRT; 365 AA.
AC Q09160;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE HLA class I histocompatibility antigen, AW-80 (A-1) alpha chain
DE precursor.
GN HLA-A OR HLA-A.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN 1;
RP SEQUENCE FROM N.A.
RX MEDLINE=94245293; PubMed=8188325;
RA Balas A., Garcia-Sanchez F., Gomez-Reino F., Vicario J.L.;
RT "Characterization of a new and highly distinguishable HLA-A allele in
RT a Spanish family.";
RL Immunogenetics 39:452-452 (1994).
RN 2;
RP SEQUENCE FROM N.A.
RA Domena J.D.;
RL Submitted (JUN-1993) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
CC -1- POLYMORPHISM: THE ONLY ALLELE OF AW-80 KNOWN IS A*8001 WHICH IS

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DR EMBL; U03754; AAC04322.1; -

DR EMBL; L18898; AAA17012.1; -

DR PIR; I59638; I38439.

DR HSP; Q95352; IHHK.

DR MIM; 142800; -

DR GO; GO:0005887; C: integral to plasma membrane; NAS.

DR GO; GO:0030106; F: MHC class I receptor activity; NAS.

DR GO; GO:0006955; P: immune response; NAS.

DR InterPro; IPRO07110; Ig-like.

DR InterPro; IPRO03597; Ig-cl.

DR InterPro; IPRO03006; Ig_MHC.

DR InterPro; IPRO01039; MHC_I.

DR Pfam; PF00047; Ig; 1.

DR Pfam; PF00129; MHC_I; 1.

DR PRINTS; PR01638; MHCCLASSI.

DR PRODOM; PD000050; MHC_I; 1.

DR SMART; SM00407; IGL1; 1.

DR PROSITE; PS00835; IG_LIKE; 1.

DR PROSITE; PS00290; IG_MHC; 1.

KW MHC I; Transmembrane; Glycoprotein; Signal.

FT SIGNAL 1 24

FT CHAIN 25 365 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN.

FT DOMAIN 25 114 AW-80(A-1) ALPHA CHAIN.

FT DOMAIN 115 206 EXTRACELLULAR ALPHA-1.

FT DOMAIN 207 298 EXTRACELLULAR ALPHA-2.

FT DOMAIN 299 308 EXTRACELLULAR ALPHA-3.

FT TRANSMEM 309 332 CONNECTING PEPTIDE.

FT DOMAIN 333 365 CYTOPLASMIC TAIL.

FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (BY SIMILARITY).

FT DISULFID 125 188 BY SIMILARITY.

FT DISULFID 227 283 BY SIMILARITY.

SQ SEQUENCE 365 AA; 40791 MW; CE1BC1CD60CA8FA8 CRC64;

Query Match 33.6%; Score 510; DB 1; Length 365;

Best Local Similarity 38.3%; Pred. No. 1.5e-35;

Matches 106; Conservative 53; Mismatches 110; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDGLSLFELGVDDQLFVYDDE--SRVPRTPWSSRISSQW 62

DB 26 SHSMRYFFTSVSRPGRGEPRFIAVGVDSDSQFVQFSDAASQRMPEAPWIEQE-EPEYW 84

QY 63 LQLSQSLKGDHMTFVDFWTFIMENHNSKE-SHTLQVILGCENQEDNS-TEGYWKYGYDG 120

DB 85 DEETRNVKAHSQNRNANLGLRGYYNQSDGSHTIQIMYGCVDGSGRFLRGYRDAYDG 144

QY 121 QDHLFPCPTDLWRRAEPRAPWTKLEWRKIRARONRAYLRDCPAQLQQLLELGRGVL 180

DB 145 KDVIALLNEDLRSTADMAAQITRKWEAR-RAEQIRAYLEGECVDGLRRLYENKETL 203

QY 181 DQOVPLPVKVTTH-VTSSVTTLCRALNYPQNTMKWLKDKQPMDAKFEFKDVLNPGD 239

DB 204 QRTDPPKTHMTHHPISDHEATLRCWLSFPABETILTQWRDGD-QTQDTLVELTRPAGD 262

QY 240 GTYQGITLAVPGEQRQYTCVQEHPLQDPLVIVE 276

DB 263 GTFQKAAVVPVGKSKRYTCHVQHEGLPEPLTRWE 299

RESULT 15

1A31_HUMAN

ID 1A31_HUMAN STANDARD; PRT; 365 AA.

AC F16189; Q98137; Q9TQ24;

DT 01-APR-1990 (Rel. 14, Created)

DT 01-APR-1993 (Rel. 25, Last sequence update)

DT 28-FEB-2003 (Rel. 41, Last annotation update)

DE HLA class I histocompatibility antigen, A-31 alpha chain precursor (Aw-19).

DE HLA-A OR HLA-A.

GN Homo sapiens (Human).

OS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

OX NCBI_TaxID=9606;

RN [1] SEQUENCE FROM N.A. (A*3101).

RX MEDLINE=90038496; PubMed=2478623;

RA Kato K., Trapani J.A., Allopenna J., Dupont B., Yang S.Y.; "Molecular analysis of the serologically defined HLA-Aw19 antigens. A genetically distinct family of HLA-A antigens comprising A29, A31, A32, and Aw33, but probably not A30.";

RT J. Immunol. 143:3371-3378(1989).

RN [2] SEQUENCE FROM N.A. (A*3101).

RX MEDLINE=92269955; PubMed=1317015;

RA Belich M.P., Madrigal J.A., Hildebrand W.H., Zemmour J., Williams R.C., Luz R., Petzl-Erler M.L., Parham P.; "Unusual HLA-B alleles in two tribes of Brazilian Indians.";

RT Nature 357:326-329(1992).

RN [3] SEQUENCE FROM N.A. (A*3101).

RX MEDLINE=96387675; PubMed=8795145;

RA Arnett K.L., Adams E.J., Parham P.; "On the sequence of A*3101.";

RT Tissue Antigens 47:428-430(1996).

RN [4] SEQUENCE OF 9-365 FROM N.A. (A*3101).

RX MEDLINE=92269956; PubMed=1589035;

RA Watkins D.I., McAdam S.N., Liu X., Stang C.R., Milford E.L., Levine C.G., Garber T.L., Dogan A.L., Lord C.I., Ghim S.H., Troup G.M., Hughes A.L., Letvin N.L.; "New recombinant HLA-B alleles in a tribe of South American Amerindians indicate rapid evolution of MHC class I loci.";

RT Nature 357:329-333(1992).

RN [5] SEQUENCE FROM N.A. (A*3104).

RA Bettinotti M.P., Dhillion G., Hackett J., Simonis T.B., Marincola F.M.; "A New HLA-A*31 allele.";

RT Submitted (MAY-1999) to the EMBL/GenBank/DBJ databases.

RN [6] SEQUENCE OF 26-206 FROM N.A. (A*3104).

RA Mitsushige Y.; "New HLA-A31 allele identified in African American population.";

RT Submitted (NOV-1998) to the EMBL/GenBank/DBJ databases.

CC -!- FUNCTION: Involved in the presentation of foreign antigens to the immune system.

CC -!- SUBUNIT: Dimer of alpha chain and a beta chain (beta-2-microglobulin).

CC -!- POLYMORPHISM: THE FOLLOWING ALLELES OF A-31 ARE KNOWN: A*3101 AND A*3104. THE SEQUENCE SHOWN IS THAT OF A*3101.

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CC -----

DR EMBL; M30578; AAA59613.1; -

DR EMBL; M84375; AAA59599.1; -

DR EMBL; L78918; AAB05976.1; -

DR EMBL; AF148863; AAD39981.1; -

DR EMBL; AF105028; AAC79721.1; -

DR EMBL; AF105027; AAC79721.1; JOINED.

DR FIR; I72170; I72170.

DR HSP; O19673; 1HSB.

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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:05:29 ; Search time 32 Seconds
(without alignments)

2225.704 Million cell updates/sec

Title: US-10-092-404-2

Perfect score: 1520

Sequence: 1 RLRSLSLHFLMGASEQDL.....RYTCQVHPGLDQPLIVME 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL_23:*
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mmc:*
8: sp_organelle:*
9: sp_phase:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*
15: sp_rvirus:*
16: sp_bacteriap:*
17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1140	75.0	358	11 Q8C2A6	Q8C2A6 mus musculus
2	1140	75.0	359	11 Q9D754	Q9D754 mus musculus
3	802	52.8	272	11 Q9R105	Q9R105 rattus norv
4	592	38.9	116	4 Q9HC69	Q9HC69 homo sapien
5	547.5	36.0	359	7 Q8HX81	Q8HX81 ornithorhyn
6	543.5	35.8	340	7 Q9BD50	Q9BD50 pongo pygma
7	542.5	35.7	334	7 Q9TOK3	Q9TOK3 homo sapien
8	542.5	35.7	341	4 Q9NPL2	Q9NPL2 homo sapien
9	542.5	35.7	341	7 Q9S460	Q9S460 homo sapien
10	542.5	35.7	341	7 Q9BCU3	Q9BCU3 pan troglod
11	540.5	35.6	354	7 Q9SHB3	Q9SHB3 anas platyr
12	539.5	35.5	341	7 Q9BCU4	Q9BCU4 pan troglod
13	530	34.9	105	4 Q9HC71	Q9HC71 homo sapien
14	521	34.3	356	7 Q8HX66	Q8HX66 sus scrofa
15	520	34.2	332	7 Q30990	Q30990 pan troglod
16	520	34.2	365	7 Q9TLP7	Q9TLP7 pan troglod

ALIGNMENTS

RESULT 1

Q8C2A6 ID Q8C2A6 PRELIMINARY; PRT; 358 AA.

AC Q8C2A6; DT 01-MAR-2003 (TrEMBLrel. 23, Created)

DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)

DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)

DB Hemochromatosis.

OS Mus musculus (Mouse).

OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.

OX NCBI_TaxID=10090;

RN [1]

RP SEQUENCE FROM N.A.

RC STRAIN=NOD; TISSUE=Thymus;

RX MEDLINE=22354683; PubMed=12466851;

RA The FANTOM Consortium.

RA the RIKEN Genome Exploration Research Group Phase I & II Team;

RT "Analysis of the mouse transcriptome based on functional annotation of

RT 60,770 full-length cDNAs."

RL Nature 420:563-573(2002).

DR EMBL; AK089886; BAC40688.1; --

SQ SEQUENCE 358 AA; 40421 MW; EE88FB6E5AAC844D CRC64;

Query Match 75.0%; Score 1140; DB 11; Length 358;

Best Local Similarity 72.2%; Pred. No. 2.5e-99;

Matches 203; Conservative 30; Mismatches 40; Indels 8; Gaps 1;

QY 4 RSHSLHFLMGASEQDLGLSFEALGYVDDQLFVFYDDSSRRVPTPWSSRISSQWL 63

29 RSHSLHFLMGASEQDLGLSFEALGYVDDQLFVFYDDSSRRVPTPWSSRISSQWL 88

64 QLSQSLKGDHMTFTVDFTWIMENHNSK-----ESHTLQVILGCMEQDNSTGYWK 115

89 HLSQSLKGDHMTFTVDFTWIMENHNSK-----ESHTLQVILGCMEQDNSTGYWK 148

116 YGVDGDHLEFCPDTLDWRAEFPRAWPTKLEWERHKIRAKONRAYLERCPQLQLEL 175

149 YGVDGDHLEFCPDTLDWRAEFPRAWPTKLEWERHKIRAKONRAYLERCPQLQLEL 208

Q9TQ66 homo sapien
Q9TQ67 homo sapien
Q8MHT1 sus scrofa
O19243 sus scrofa
Q9MYG4 homo sapien
Q29747 homo sapien
Q9S196 homo sapien
Q8SPA9 sus scrofa
Q8HX63 sus scrofa
Q8HX61 sus scrofa
Q9MYI5 homo sapien
Q19356 macaca mula
O02944 macaca mula
Q98030 papio anubi
Q98031 papio anubi
Q9UK37 homo sapien
O02947 macaca mula
O02946 macaca mula
O19756 homo sapien
O02945 macaca mula
Q30886 pan paniscu
Q9MXI5 pan troglod
Q9MWK4 gorilla gor
Q9MXI6 pan troglod
Q9MXM7 pan troglod
Q9GJ24 homo sapien
Q8SPA4 sus scrofa
Q30900 pan paniscu
Q9BCN0 homo sapien

```

QY 176 GRGVLQVQVPLVKVTHVTSVTLRCALNYYPQNTMKLQKQPMDAKEFEFKDVL 235
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 209 GRGVLQVQVPLVKVTHVTSVTLRCALNYYPQNTMKLQKQPMDAKEFEFKDVL 268
QY 236 PNGDGTQYQWITLAVPPGEQRYTCQVEHPGLDQPLVIWE 276
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 269 PNGDGTQYQWITLAVPPGEQRYTCQVEHPGLDQPLVIWE 309

RESULT 2
Q9D754 PRELIMINARY; PRT; 359 AA.
AC Q9D754;
DT 01-JUN-2001 (T-EMBLrel. 17, Created)
DT 01-JUN-2001 (T-EMBLrel. 17, Last sequence update)
DT 01-MAR-2003 (T-EMBLrel. 23, Last annotation update)
DE Adult male tongue cDNA, RIKEN full-length enriched library,
DE clone:2310032M04, full insert sequence.
GN HFE.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RC SEQUENCE FROM N.A.
RX MEDLINE=2108560; PubMed=11217851;
RA Kawai J., Shingawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
RA Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,
RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
RA Fleischmann W., Gaasterland T., Giesi C., King B., Kochiwa H.,
RA Kuehl P., Lewis S., Matsuo Y., Nikaide I., Pesole G., Quackenbush J.,
RA Schram L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
RA Guncicich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Suzuki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
RA Suzuki H., Toyo-oka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
RA Wynshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,
RA Havaehizaki Y.
RT "Functional annotation of a full-length mouse cDNA collection.";
RL Nature 409:685-690(2001).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
CC ENBL; AK009581; BAB26373.1; -.
DR HSSP; Q30201; 1A6Z.
DR MGD; MGI:109191; Hfe.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR PROSITE; PS00290; IG_MHC; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS00835; IG-LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
SQ SEQUENCE 359 AA; 40534 MW; 586657B7F9FF20B4 CRC64;

Query Match 75.0%; Score 1140; DB 11; Length 359;
Best Local Similarity 72.2%; Pred. No. 2.5e-99;
Matches 203; Conservative 30; Mismatches 40; Indels 8; Gaps 1;

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QY 4 RSHSLHYLPMGASEQDLGLSFEALGYVDDQLFVYVDDSRVRPRTPWVSRRISQMWL 63
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Db 30 RSHSLHYLPMGASEQDLGLSFEALGYVDDQLFVYVDDSRVRPRTPWVSRRISQMWL 89
QY 64 QLSQSLKGDHMTFTVDFTIMENHNHSHK-----ESHTLOVILGCEMOEDNSTGYWK 115
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 90 HUSQSLKGDHMTFTVDFTIMGNHNSKVTKLGVVSESHILOVILGCEVHEDNSTSGFWR 149
QY 116 YGYDGDHLEFCFCDTLDWRAAPRAWPTKLEWHERKIRARQNRVYLERDCPAQLQELLE 175
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 150 YGYDGDHLEFCFCKTLNWSAEPGAWATKVEWDEKIRAKQNRDYLEKDCPEQLKELLE 209
QY 176 GRGVLQVQVPLVKVTHVTSVTLRCALNYYPQNTMKLQKQPMDAKEFEFKDVL 235
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 210 GRGVLQVQVPLVKVTHVTSVTLRCALNYYPQNTMKLQKQPMDAKEFEFKDVL 269
QY 236 PNGDGTQYQWITLAVPPGEQRYTCQVEHPGLDQPLVIWE 276
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 270 PNGDGTQYQWITLAVPPGEQRYTCQVEHPGLDQPLVIWE 310

RESULT 3
Q9RI05 PRELIMINARY; PRT; 272 AA.
AC Q9RI05;
DT 01-MAY-2000 (T-EMBLrel. 13, Created)
DT 01-MAY-2000 (T-EMBLrel. 13, Last sequence update)
DT 01-MAR-2003 (T-EMBLrel. 23, Last annotation update)
DE Hemochromatosis gene product HFE splice variant delE2.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RC SEQUENCE FROM N.A.
RX STRAIN=Wistar; TISSUE=Testis;
RA Liew Y.-F., Shaw N.-S.;
RT "Alternative splice variant of the hemochromatosis gene HFE in iron
RT overloaded rats.";
RL Submitted (AUG-1999) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
CC ENBL; AF176534; AAD49965.1; -.
DR HSSP; Q30201; 1A6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS00835; IG-LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
SQ SEQUENCE 272 AA; 30757 MW; 1D91063CCBEF5502 CRC64;

Query Match 52.8%; Score 802; DB 11; Length 272;
Best Local Similarity 75.1%; Pred. No. 1.4e-67;
Matches 139; Conservative 22; Mismatches 24; Indels 0; Gaps 0;

QY 92 ESHTLOVILGCEMOEDNSTGYWKYDGDHLEFCFCDTLDWRAAPRAWPTKLEWHERK 151
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 39 ESHTLOVILGCEVHEDNSTGSGFWKYGVDGDHLEFCFCKTLNWSAEPGAWATKVEWHER 98
QY 152 IRARQNRVYLERDCPAQLQELLEGRVLDQVPPPLVKVTHVTSVTLRCALNYYPQ 211
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 99 IRARQNRVYLERDCPAQLQELLEGRVLDQVPPPLVKVTHVTSVTLRCALNYYPQ 158
QY 212 NITMKWLKQPMDAKEFEFKDVL PNGDGTQYQWITLAVPPGEQRYTCQVEHPGLDQPL 271

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Db 83 RYTQLLRGQOMPFKVELKRLQRHYNHS -GSHTYQRMIGICELLBGSGTTGFLQVAYDQDF 141
Qy 124 LEFCPTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 142 LIFNKDTLSLWADVNVHTTIKRAWEANOHELOQKQWLEECIAWLKRFLEYKDTLQRT 201
Qy 184 VPPLVKVTHVHT-SSVTTLCRALNYYPNITMKWLKDKQPMDAKEFFPKDVLPLNGDGT 242
Db 202 EPLVRVNRKETFPFGVTTLCFAHGFYPEIYMTWMKGEEI-VQEMDYDGLILPSGDGT 260
Qy 243 QGWITLAVPPGEEQRYTCQVEHPGLDQPLIV 273
Db 261 QTWASFELDPQSSNLSYCHVEHCGVHMVLQV 291

RESULT 7

Q9TK3 ID Q9TK3 PRELIMINARY; PRT; 334 AA.
AC Q9TK3
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE MHC class I-related protein MRL (Fragment).
GN MRL.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Placenta;
RX MEDLINE=99003494; PubMed=9784382;
RA Yamaguchi H., Kurosawa Y., Hashimoto K.;
RT "Expanded genomic organization of conserved mammalian MHC class I-
RT related genes, human MRL and its murine ortholog.";
RL Biochem. Biophys. Res. Commun. 250:558-564 (1998).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; AF073485; AAC72900.1; JOINED.
DR EMBL; AF073484; AAC72900.1; JOINED.
DR HSSP; Q30201; IA6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR ProDom; PD000050; MHC I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG-LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
FT NON TER 1
SQ SEQUENCE 334 AA; 38586 MW; 4C3E8A248A39A4 CRC64;

Query Match 35.7%; Score 542.5; DB 7; Length 334;
Best Local Similarity 39.5%; Pred. No. 6e-43;
Matches 107; Conservative 50; Mismatches 111; Indels 3; Gaps 3;
Qy 4 RSHSLYLFMGASEQDLGLSLFALGVYDDQLFVYDDSRREPRTPWSSRISSQMWL 63
Db 16 RTHSLRYFLGVSDPIHGVPFISGVYDHPITTYDSVTRQKEPRAPMAENLADPHE 75
Qy 64 QLSQSLKGWDMFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDQDH 123
Db 76 RYTQLLRGQOMPFKVELKRLQRHYNHS -GSHTYQRMIGICELLBGSGTTGFLQVAYDQDF 134
Qy 124 LEFCPTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 135 LIFNKDTLSLWADVNVHTTIKRAWEANOHELOQKQWLEECIAWLKRFLEYKDTLQRT 194

Qy 184 VPPLVKVTHVHT-SSVTTLCRALNYYPNITMKWLKDKQPMDAKEFFPKDVLPLNGDGT 242
Db 195 EPLVRVNRKETFPFGVTTLCFAHGFYPEIYMTWMKGEEI-VQEMDYDGLILPSGDGT 253
Qy 243 QGWITLAVPPGEEQRYTCQVEHPGLDQPLIV 273
Db 254 QWASIELDPQSSNLSYCHVEHCGVHMVLQV 284

RESULT 8

Q9NPL2 ID Q9NPL2 PRELIMINARY; PRT; 341 AA.
AC Q9NPL2
DT 01-OCT-2000 (TREMBlrel. 15, Created)
DT 01-OCT-2000 (TREMBlrel. 15, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE MRL protein.
GN MRL.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Peripheral blood;
RX MEDLINE=20470599; PubMed=11019920;
RA Parra-Cuadrado J.F., Navarro P., Mirones I., Setien F., Oteo M.,
RA Martinez-Naves E.;
RT "A study on the polymorphism of human MHC class I-related MRL gene and
RT identification of an MRL-like pseudogene.";
RL Tissue Antigens 56:170-172 (2000).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; AJ249778; CAB77667.1; -.
DR HSSP; Q30201; IA6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR ProDom; PD000050; MHC I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG-LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
SQ SEQUENCE 341 AA; 39366 MW; 2990C1F3F0A1CAD9 CRC64;

Query Match 35.7%; Score 542.5; DB 4; Length 341;
Best Local Similarity 39.5%; Pred. No. 6.2e-43;
Matches 107; Conservative 50; Mismatches 111; Indels 3; Gaps 3;
Qy 4 RSHSLYLFMGASEQDLGLSLFALGVYDDQLFVYDDSRREPRTPWSSRISSQMWL 63
Db 23 RTHSLRYFLGVSDPIHGVPFISGVYDHPITTYDSVTRQKEPRAPMAENLADPHE 82
Qy 64 QLSQSLKGWDMFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDQDH 123
Db 83 RYTQLLRGQOMPFKVELKRLQRHYNHS -GSHTYQRMIGICELLBGSGTTGFLQVAYDQDF 141
Qy 124 LEFCPTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 142 LIFNKDTLSLWADVNVHTTIKRAWEANOHELOQKQWLEECIAWLKRFLEYKDTLQRT 201
Qy 184 VPPLVKVTHVHT-SSVTTLCRALNYYPNITMKWLKDKQPMDAKEFFPKDVLPLNGDGT 242
Db 202 EPLVRVNRKETFPFGVTTLCFAHGFYPEIYMTWMKGEEI-VQEMDYDGLILPSGDGT 260
Qy 243 QGWITLAVPPGEEQRYTCQVEHPGLDQPLIV 273

44.


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Matches 94; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
QY 89 HSKESHTLOVILGCEMEDNSTEGYKYGVDGQDHLFCFDTLDWRAAEPRAPMTKLEWE 148
DB 1 HTKESHTLOVILGCEMEDNSTEGYKYGVDGQDHLFCFDTLDWRAAEPRAPMTKLEWE 60
QY 149 RHKIRARONRAYLERDCPAQLQQLLELGRGVLDQVPPV 188
DB 61 GHKVRARQAYLERDCPAQLQQLLELGRGVLDQVPPV 100

RESULT 14
Q8HX66 PRELIMINARY; PRT; 356 AA.
AC Q8HX66
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE MHC class I antigen (Fragment).
GN SLA-1.
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
NCBI_TaxID=9823;
RN [1]
RP SEQUENCE FROM N.A.
RA Martens G.W., Baker J.E., Smith D.M.;
RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY135589; AN35107.1; -.
FT NON TER 1
SQ SEQUENCE 356 AA; 3585 MW; 94FC7A461DBF555B CRC64;

Query Match 34.3%; Score 521; DB 7; Length 356;
Best Local Similarity 39.9%; Pred. No. 7e-41;
Matches 110; Conservative 48; Mismatches 110; Indels 8; Gaps 7;
QY 6 HSLHYLFMGASEODLGLSLFEALGYVDDQLFVYDDE--SRVPEPTPWSSRISSQMWL 63
DB 19 HSLRYFYTAVRPDLGDSRFIAVGYDDTQVFRFSDAPNPRMEPRAPMTQOE--GOEYWD 77
QY 64 QLSQSLKGWDHMTVDFTWIMENHNHKSKE--SHTLQVILGCEMEDN--STEGYKYGVDGQ 121
DB 78 EETRNANGSQNDVRDLKTLRGYNSQSEAGSHIIQRMYGCDVPGDGLLGRYDQDAYDGA 137
QY 122 DHLEFCFDTLDWRAAEPRAPMTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVLD 181
DB 138 DYIALNEDLRSWTAADTAQAQITKRKWEAANV-AEQERSYLEGTCVWELQKYLEMGKDTLQ 196
QY 182 QVPPPLVKVTHHTSSV--TTLRCALNYYPONTIMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 197 RAEPKTHVTRHPSSDLGVTLRCWALGFYFKEISLTWQREGQD--QSQDMELVETRPSGDG 255
QY 241 TYOGWITLAVPPGGEORYTCQVHPGLDQPLVIWE 276
DB 256 TFQKWAALVPPGGEQSYTCHVQHEGLPKPLTLRWD 291

RESULT 15
Q30990 PRELIMINARY; PRT; 332 AA.
AC Q30990
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Chimpanzee MHC class I Chla chain (Fragment).
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=89235215; PubMed=2715640;
RA Farham P., Lawlor D.A., Lomen C.E., Ennis P.D.;
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RT Diversity and diversification of HLA-A,B,C alleles.";
RL J. Immunol. 142:3937-3950(1989).
CC -I- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -I- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; M24047; AAA35426.1; -.
DR HSSP; Q95352; IHHK.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_c1.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IgC1; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
FT NON TER 332
SQ SEQUENCE 332 AA; 37433 MW; 9AA9A55DF9E79360 CRC64;

Query Match 34.2%; Score 520; DB 7; Length 332;
Best Local Similarity 40.1%; Pred. No. 7.9e-41;
Matches 111; Conservative 44; Mismatches 114; Indels 8; Gaps 7;
QY 5 HSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDE--SRVPEPTPWSSRISSQMW 62
DB 26 SHSMRYFTTSVSRPGEGEPRFIAVGYDDTQVFRFSDAASQRMPEPRAPWIOQE--GPEYW 84
QY 63 LQLSOSLKGWDHMTVDFTWIMENHNHKSKE--SHTLQVILGCEMEDNS--TEGYKYGVDG 120
DB 85 DQETRSAAHSQTDVRDLGTLRGYNSQSEDSHTTIQIMYGCDVSGDGRFLRGYRQDAYDG 144
QY 121 QHLEFCFDTLDWRAAEPRAPMTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVLD 180
DB 145 KYDIALNEDLRSWTAADMAAQITKRKWEAAH--AAEQRAYLEGTCVWELRLRYLENGKETL 203
QY 181 DOQVPLVKVTHH--VTSSVTLRCALNYYPONTIMKWLKDKQPMDAKEPEPKDVLNPGD 239
DB 204 QRTDPPKTHMTHTPISDHEATLRCWALGFYFPAEITLTWQDGED--QTQDTLVELTRPADG 262
QY 240 GYQGWITLAVPPGGEORYTCQVHPGLDQPLVIWE 276
DB 263 GTFQKWAALVPPGGEQRYTCHVQHEGLPKPLTLRWE 299

Search completed: August 5, 2003, 13:10:00
Job time : 34 secs
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:10:04 ; Search time 33 Seconds
(without alignments)
993.264 Million cell updates/sec

Title: US-10-092-404-2
Perfect score: 1520
Sequence: 1 RLRSLSLHLYFMGASQDL.....RYTCQVHEGLDQPLIVIME 276

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 451899 seqs, 118759770 residues

Total number of hits satisfying chosen parameters: 451899

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :				Published Applications AA:	
1:	/cgn2_6/ptodata/1/pubpaa/US07_PUBCOMB.pep.*				
2:	/cgn2_6/ptodata/1/pubpaa/PCT_NEW_PUB.pep.*				
3:	/cgn2_6/ptodata/1/pubpaa/US06_NEW_PUB.pep.*				
4:	/cgn2_6/ptodata/1/pubpaa/US06_PUBCOMB.pep.*				
5:	/cgn2_6/ptodata/1/pubpaa/US07_NEW_PUB.pep.*				
6:	/cgn2_6/ptodata/1/pubpaa/PCTUS_PUBCOMB.pep.*				
7:	/cgn2_6/ptodata/1/pubpaa/US08_NEW_PUB.pep.*				
8:	/cgn2_6/ptodata/1/pubpaa/US08_PUBCOMB.pep.*				
9:	/cgn2_6/ptodata/1/pubpaa/US09A_PUBCOMB.pep.*				
10:	/cgn2_6/ptodata/1/pubpaa/US09B_PUBCOMB.pep.*				
11:	/cgn2_6/ptodata/1/pubpaa/US09C_PUBCOMB.pep.*				
12:	/cgn2_6/ptodata/1/pubpaa/US09_NEW_PUB.pep.*				
13:	/cgn2_6/ptodata/1/pubpaa/US10A_PUBCOMB.pep.*				
14:	/cgn2_6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*				
15:	/cgn2_6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*				
16:	/cgn2_6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*				
17:	/cgn2_6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*				
18:	/cgn2_6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*				

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1520	100.0	276	15	US-10-092-404-2
2	1513	99.5	276	15	US-10-092-404-1
3	1513	99.5	348	12	US-09-981-606-1
4	1493	98.2	276	15	US-10-092-404-3
5	514	33.8	92	14	US-10-016-634A-120
6	506	33.3	280	15	US-10-073-300-6
7	506	33.3	415	15	US-10-073-300-5
8	492	32.4	298	15	US-10-205-823-40
9	492	32.4	298	15	US-10-205-823-42
10	492	32.4	298	15	US-10-177-293-23
11	477	31.4	542	15	US-10-015-535-32
12	477	31.4	542	15	US-10-015-535-34
13	475	31.2	542	15	US-10-015-535-36
14	474	31.2	540	15	US-10-015-535-22
15	474	31.2	541	15	US-10-015-535-28

16	474	31.2	542	15	US-10-015-535-24	Sequence 24, Appl
17	474	31.2	542	15	US-10-015-535-26	Sequence 26, Appl
18	448	29.5	332	9	US-09-870-521-3	Sequence 3, Appli
19	445	29.3	540	15	US-10-015-535-30	Sequence 30, Appli
20	444	29.2	334	9	US-09-870-521-4	Sequence 4, Appli
21	358.5	23.6	170	9	US-09-925-301-1307	Sequence 1307, Ap
22	336	22.1	271	9	US-09-925-301-1431	Sequence 1431, Ap
23	279	18.4	181	11	US-09-013-077A-13	Sequence 13, Appl
24	275	18.1	145	9	US-09-810-560-8	Sequence 8, Appli
25	243	16.0	184	10	US-09-858-580-21	Sequence 21, Appl
26	243	16.0	184	11	US-09-847-172-21	Sequence 21, Appl
27	226	14.9	91	9	US-09-864-761-38005	Sequence 38005, A
28	223.5	14.7	171	15	US-10-144-929-116	Sequence 116, App
29	223	14.7	91	9	US-09-864-761-35461	Sequence 35461, A
30	210.5	13.8	104	9	US-09-925-302-835	Sequence 835, App
31	208.5	13.7	183	15	US-10-036-542-62	Sequence 62, Appl
32	207	13.6	117	9	US-09-810-560-9	Sequence 9, Appli
33	196.5	12.9	93	9	US-09-864-761-39479	Sequence 39479, A
34	196.5	12.9	110	9	US-09-864-761-35339	Sequence 35339, A
35	196.5	12.9	114	9	US-09-864-761-37988	Sequence 37988, A
36	174.5	11.5	261	10	US-09-925-664-30	Sequence 30, Appl
37	174	11.4	411	14	US-10-015-536-17	Sequence 17, Appl
38	173	11.4	110	10	US-09-796-692-799	Sequence 799, App
39	173	11.4	110	10	US-09-796-692-2139	Sequence 2139, Ap
40	173	11.4	110	15	US-10-040-862-799	Sequence 799, App
41	173	11.4	110	15	US-10-040-862-2139	Sequence 2139, Ap
42	171.5	11.3	285	10	US-09-756-983-24	Sequence 24, Appl
43	167	11.0	772	9	US-09-815-837-74	Sequence 74, Appl
44	166.5	11.0	448	14	US-10-081-281-111	Sequence 111, App
45	166	10.9	246	9	US-09-989-722-225	Sequence 225, App

ALIGNMENTS

RESULT 1

US-10-092-404-2
; Sequence 2, Application US/10092404
; Publication No. US20030073627A1
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; Bjorkman, Pamela J.
; Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; AND IRON DEFICIENCY DISEASES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/10/092,404
; FILING DATE: 04-Mar-2002
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:

TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 2:
US-10-092-404-2

Query Match 100.0%; Score 1520; DB 15; Length 276;
Best Local Similarity 100.0%; Pred. No. 1.7e-146;
Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWSSRISSQ 60
QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQLSQSLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDHLFCFPTDLWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQQLLELGRGVL 180
DB 121 QDHLFCFPTDLWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQQLLELGRGVL 180
QY 181 DQVPLVAVKTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEKDVLPNGDG 240
DB 181 DQVPLVAVKTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEKDVLPNGDG 240
QY 241 TYQGWITLAVPGEERQYTCQVEHPGLDQPLIWIWE 276
DB 241 TYQGWITLAVPGEERQYTCQVEHPGLDQPLIWIWE 276

RESULT 2

US-10-092-404-1
Sequence 1, Application US/10092404
Publication No. US20030073627A1
GENERAL INFORMATION:
APPLICANT: Feder, John N.
Bjorkman, Pamela J.
Schatzman, Randall C.
TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
AND IRON DEFICIENCY DISEASES
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds, LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows
SOFTWARE: FastSeq for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/10/092,404
FILING DATE: 04-Mar-2002
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/09/094,964
FILING DATE: June 12, 1998
APPLICATION NUMBER: 08/876,010
FILING DATE: June 13, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M

REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0074-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-10-092-404-1

Query Match 99.5%; Score 1513; DB 15; Length 276;
Best Local Similarity 99.6%; Pred. No. 8.6e-146;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWSSRISSQ 60
QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQLSQSLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDHLFCFPTDLWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQQLLELGRGVL 180
DB 121 QDHLFCFPTDLWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQQLLELGRGVL 180
QY 181 DQVPLVAVKTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEKDVLPNGDG 240
DB 181 DQVPLVAVKTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEKDVLPNGDG 240
QY 241 TYQGWITLAVPGEERQYTCQVEHPGLDQPLIWIWE 276
DB 241 TYQGWITLAVPGEERQYTCQVEHPGLDQPLIWIWE 276

RESULT 3

US-09-981-606-2
Sequence 2, Application US/09981606
Publication No. US20030129595A1
GENERAL INFORMATION:
APPLICANT: Rothenberg et al.
TITLE OF INVENTION: Mutations associated with iron disorders
FILE REFERENCE: 24065-004CON
CURRENT APPLICATION NUMBER: US/09/981,606
CURRENT FILING DATE: 2002-10-16
PRIOR APPLICATION NUMBER: 09/277,457
PRIOR FILING DATE: 1999-03-26
NUMBER OF SEQ ID NOS: 30
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 2
LENGTH: 348
TYPE: PRT
ORGANISM: Homo sapiens
US-09-981-606-2

Query Match 99.5%; Score 1513; DB 12; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.2e-145;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142

QY 121 QHLEFCPTDLWRAAEPRAMPKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 180
Db 143 QHLEFCPTDLWRAAEPRAMPKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
Db 203 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPPGEORQYTCQVEHPGLDQPLIWIWE 276
Db 263 TYQGWITLAVPPGEORQYTCQVEHPGLDQPLIWIWE 298
RESULT 4
US-10-092-404-3
; Sequence 3, Application US/10092404
; Publication No. US20030073627A1
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; Bjorkman, Pamela J.
; Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; AND IRON DEFICIENCY DISEASES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/10/092,404
; FILING DATE: 04-Mar-2002
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 276 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; SEQUENCE DESCRIPTION: SEQ ID NO: 3:
US-10-092-404-3
Query Match 98.2%; Score 1493; DB 15; Length 276;
Best Local Similarity 98.9%; Pred. No. 9.3e-144;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDYDDESRVRPPTPWVSSRISQ 60
Db 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDYDDESRVRPPTPWVSSRISQ 60
QY 61 MWLQLSLSKGDHMTFTVDFTIMNHNHNSKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120

Db 61 MWLQLSLSKGDHMTFTVDFTIMNHNHNSKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QHLEFCPTDLWRAAEPRAMPKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 180
Db 121 QHLEFCPTDLWRAAEPRAMPKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 180
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
Db 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
QY 241 TYQGWITLAVPPGEORQYTCQVEHPGLDQPLIWIWE 276
Db 241 TYQGWITLAVPPGEORQYTCQVEHPGLDQPLIWIWE 276
RESULT 5
US-10-016-634A-120
; Sequence 120, Application US/10016634A
; Publication No. US20020192666A1
; GENERAL INFORMATION:
; APPLICANT: Sun, Yongming
; APPLICANT: Recipon, Herve
; APPLICANT: Ghosh, Malavika
; APPLICANT: Liu, Chenghua
; TITLE OF INVENTION: Compositions and Methods Relating to Colon Specific Genes and Proteins
; FILE REFERENCE: DEX-0255
; CURRENT APPLICATION NUMBER: US/10/016,634A
; CURRENT FILING DATE: 2001-10-31
; PRIOR APPLICATION NUMBER: US 60/244,258
; PRIOR FILING DATE: 2000-10-31
; NUMBER OF SEQ ID NOS: 176
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 120
; LENGTH: 92
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-016-634A-120
Query Match 33.8%; Score 514; DB 14; Length 92;
Best Local Similarity 100.0%; Pred. No. 9.5e-45;
Matches 92; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 92 ESHTLQVILGCEMQEDNSTEGYWKYGYDGQDHLEFCPTDLWRAAEPRAMPKLEWERHK 151
Db 1 ESHTLQVILGCEMQEDNSTEGYWKYGYDGQDHLEFCPTDLWRAAEPRAMPKLEWERHK 60
QY 152 IRARONRAYLERDPCPAQLQQLLELGRGVLDQ 183
Db 61 IRARONRAYLERDPCPAQLQQLLELGRGVLDQ 92
RESULT 6
US-10-073-300-6
; Sequence 6, Application US/10073300
; Publication No. US20030003535A1
; GENERAL INFORMATION:
; APPLICANT: Reiter, Yoram
; TITLE OF INVENTION: SINGLE CHAIN CLASS I MAJOR HISTO- COMPATIBILITY COMPLEXES
; FILE REFERENCE: 02/23339
; CURRENT APPLICATION NUMBER: US/10/073,300
; CURRENT FILING DATE: 2002-06-25
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 6
; LENGTH: 280
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-073-300-6
Query Match 33.3%; Score 506; DB 15; Length 280;
Best Local Similarity 39.4%; Pred. No. 2.7e-43;
Matches 109; Conservative 45; Mismatches 115; Indels 8; Gaps 7;

QY 182 QQVPLVKVTHV-TSSVTLRCRALNYYPONITMKWLKDKQPMDAKBEFEPKDVLPNGDG 240
Db 234 RTDSPKAVTHHSRPEKVTLCRCWALGFYPADITLTWQNGEEL-IQDMELVETRPAGDG 292
QY 241 TYQGWITLAVPGEQRYTCQVEHFGLDQPLIVIE 276
Db 293 TFOKASVVVPLGKEQYTCVYHQGLPEPLTLRWE 328
Search completed: August 5, 2003, 13:21:55
Job time : 34 secs

QY 6 HSLHYLFMGASEODLGLSFEALGYVDDQLFVYDD--ESRRVEPRTPWSSRISSQWML 63
Db 144 HSLRYFVTAVSRPGLGEPRYMEVGYDDTEFVRPDSDAENPRYEPARMWEQE-GPEYWE 202
QY 64 QLSQSLKGMHMTVDFTIMENHNHSHK-ESHTLQVILGCEMOEDNS-TEGYWKYGYDQG 121
Db 203 RETQKAKGNEQSRVLDRLTLGGYNQSGSHTIQVISGCEVSDGRLRLGYQYAYDGC 262
QY 122 DHLEFCPDTLDWAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLOQLLELGRGVL 181
Db 263 DYIALNEDLKTWTAADMAALITKHKEQAG-EAERLRAVLEGTVCVWLRRLKNGNATLL 321
QY 182 QQVPLVKVTHV-TSSVTLRCRALNYYPONITMKWLKDKQPMDAKBEFEPKDVLPNGDG 240
Db 322 RTDSPKAVTHHSRPEKVTLCRCWALGFYPADITLTWQNGEEL-IQDMELVETRPAGDG 380
QY 241 TYQGWITLAVPGEQRYTCQVEHFGLDQPLIVIE 276
Db 381 TFOKASVVVPLGKEQYTCVYHQGLPEPLTLRWE 416

RESULT 15
US-10-015-535-28
; Sequence 28, Application US/10015535
; Publication No. US20030036506A1
; GENERAL INFORMATION:
; APPLICANT: Kranz, David M.
; APPLICANT: Brophy, Susan
; TITLE OF INVENTION: Mutated Class I Major Histocompatibility proteins and
; TITLE OF INVENTION: Complexes
; FILE REFERENCE: 100-00
; CURRENT APPLICATION NUMBER: US/10/015,535
; CURRENT FILING DATE: 2001-12-10
; PRIOR APPLICATION NUMBER: 60/254,495
; PRIOR FILING DATE: 2000-12-08
; NUMBER OF SEQ ID NOS: 37
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 28
; LENGTH: 541
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; OTHER INFORMATION: peptide
US-10-015-535-28

Query Match 31.2%; Score 474; DB 15; Length 541;
Best Local Similarity 39.5%; Pred. No. 1.2e-39;
Matches 109; Conservative 40; Mismatches 119; Indels 8; Gaps 7;
QY 6 HSLHYLFMGASEODLGLSFEALGYVDDQLFVYDD--ESRRVEPRTPWSSRISSQWML 63
Db 56 HSLRYFVTAVSRPGLGEPRYMEVGYDDTEFVRPDSDAENPRYEPARMWEQE-GPEYWE 114
QY 64 QLSQSLKGMHMTVDFTIMENHNHSHK-ESHTLQVILGCEMOEDNS-TEGYWKYGYDQG 121
Db 115 RETQKAKGNEQSRVLDRLTLGGYNQSGSHTIQVISGCEVSDGRLRLGYQYAYDGC 174
QY 122 DHLEFCPDTLDWAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLOQLLELGRGVL 181
Db 175 DYIALNEDLKTWTAADMAALITKHKEQAG-EAERLRAVLEGTVCVWLRRLKNGNATLL 233

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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:06:39 ; Search time 15 Seconds
(without alignments)
1769.504 Million cell updates/sec

Title: US-10-092-404-3
Perfect score: 1514
Sequence: 1 RLLRSLSLHLYFMGASEQDL.....RYTCQVEHPGLDQPLIVWE 276

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0
Maximum DB seq length: 2000000000
Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 76.*

1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	ID	Description
1	1129	74.6	359	2 JCS382	hereditary hemochromatosis precursor - mouse
2	530.5	35.0	341	2 A57136	class I histocomp
3	517	34.1	361	1 HLRB	MHC class I histoc
4	517	34.1	361	2 I46858	MHC class I histoc
5	514	33.9	332	2 S06424	MHC class I histoc
6	511	33.8	365	2 I36961	MHC class I histoc
7	510	33.7	361	2 B27638	MHC class I histoc
8	509	33.6	365	2 I83063	All.2 - human
9	508	33.6	365	2 A47636	MHC class I histoc
10	508	33.6	365	2 I56039	HLA-A30.3 precursor
11	506	33.4	370	1 HLHUA3	MHC class I histoc
12	504	33.3	365	2 I38439	MHC class I histoc
13	503	33.2	365	2 I37542	MHC class I histoc
14	503	33.2	365	2 I38442	gene HLA-A-0205 pr
15	503	33.2	365	2 I61902	MHC class I histoc
16	502	33.2	365	2 I72170	MHC class I histoc
17	502	33.2	365	2 I38441	gene HLA-A-6802 pr
18	501	33.1	355	2 T28149	MHC class I histoc
19	500	33.0	365	1 HLHUA2	MHC class I histoc
20	500	33.0	365	2 I37482	MHC class I histoc
21	500	33.0	365	2 I38519	MHC class I histoc
22	500	33.0	365	2 I84448	MHC class I histoc
23	499	33.0	365	2 I38610	MHC class I histoc
24	499	33.0	365	2 I37470	HLA-A*0210 - human
25	498	32.9	364	2 S03535	MHC class I histoc
26	497	32.8	365	2 I37476	MHC class I histoc
27	497	32.8	365	2 I37478	MHC class I histoc
28	497	32.8	365	2 I38443	Gene HLA-A-0203 pr
29	497	32.8	365	2 I61857	MHC HLA-A2.4a chain

MHC class I protei
MHC class I histoc
MHC class I histoc
MHC class I histoc
MHC class I histoc
MHC class I histoc
zinc-alpha 2-glyco
MHC class I histoc
class I histocompa
HLA-AW24 protein -
HLA-AW34.2 antigen
MHC class I histoc
MHC class I histoc
HLA-AW33.1, HLA-AW
MHC class I histoc
MHC class I histoc
major histocompati

ALIGNMENTS

RESULT 1

JCS382 hereditary hemochromatosis protein precursor - mouse

C;Species: Mus musculus (house mouse)

C;Date: 02-Jun-1997 #sequence_revision 18-Jul-1997 #text_change 05-Nov-1999

C;Accession: JCS382

R;Hashimoto, K.; Hirai, M.; Kurosawa, Y.

Biochem. Biophys. Res. Commun. 230, 35-39, 1997

A;Title: Identification of a mouse homolog for the human hereditary haemochromatosis ca

A;Reference number: JCS382; MUID:97148566; PMID:9020055

A;Accession: JCS382

A;Status: nucleic acid sequence not shown

A;Molecule type: DNA

A;Residues: 1-359 <HAS>

A;Cross-references: GB:U66849; NID:gl519484; PIDN:AAB07525.1; PID:gl519485

C;Comment: This protein plays a role in iron metabolism.

C;Genetics:

A;Gene: mr2

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;1-29/Domain: signal sequence #status predicted <SIG>

F;30-359/Product: hereditary haemochromatosis protein #status predicted <MAT>

F;30-117/Domain: alpha 1 #status predicted <ALF1>

F;118-217/Domain: alpha 2 #status predicted <ALF2>

F;218-309/Domain: alpha 3 #status predicted <ALF3>

F;314-340/Domain: transmembrane #status predicted <TRM>

F;341-359/Domain: intracellular #status predicted <INT>

Query Match 74.6%; Score 1129; DB 2; Length 359;

Best Local Similarity 71.9%; Pred. No. 1.9e-86;

Matches 202; Conservative 30; Mismatches 41; Indels 8; Gaps 1;

QY 4 RSHSLHLYFMGASEODLGLSFEALGYDDQLFVYDHSRVRVPTWVSRISSQWL 63

Db 30 RSHSLHLYFMGASEPDLGLPLFEARGYDDQLFVSYNHSRAEPRAPIWLEQTSQWL 89

QY 64 QLSQSLKGDHMTFTVDFTIMENHASK-----ESHTLVQILGCEMOEDNSTEGYWK 115

Db 90 HLSQSLKGDHMTFTVDFTIMGNVHNSKVTGLGVVSESHILQVLGCEVHNSSTGGFR 149

QY 116 YGYDQDLEFCPDPTLDWRAAPRAWPTYKLEWERHKIRARQNRAYLERDCAQLQLLEL 175

Db 150 YGYDQDHLFCFCKTLNNSAEPGAWATKVEWDEHKIRAKQNRDYLEKDCPEQLKLEL 209

QY 176 GGVLDQQVPLAVKTHVTSSVTTLRCALNYYPONITMKWLKQKQMDAKEPEPKDVL 235

Db 210 GRGVLGQQVPLVKTTRHWASTGTSRLRQALDFFQNTIMRWLKNQPLDKVDNPKVL 269

QY 236 PNGDGTQGWITLAVPPGEQRYTCQVEHPGLDQPLIVWE 276

Db 270 PNGDGTQGWITLAVAPGDETRFTCQVEHPGLDQPLTASWE 310

F;110/Binding site: carbohydrate (Asn) (covalent) #status predicted
F;125-188,227-283/Disulfide bonds: #status predicted

Query Match 34.1%; Score 517; DB 1; Length 361;
Best Local Similarity 40.1%; Pred. No. 1.7e-35;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7

QY 5 SHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDYHE--SRRVEPTPWSSRISSQW 62
DB SHSRYFYTSVRPGLGEPRFIIVGYDDTQVRFDSDAASPRMEQAPWM-GQVEPEYW 84
QY 63 LQLSOSLKGWDHMTVDFTWIMENHNASKB-SHTLQVILGCEMQEDNS-TGYWKYGYDG 120
DB DQQTQIAKTAQTFRVNLNTALRYNQSAAGSHTFQTMFGCEVWADGRFFHGYQYAYDG 144
QY 121 QDALEFCPDTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQLLELGRGVL 180
DB 145 ADVIALNEDLRSWTAADTAQNTQKWEAAG-EAERHRAVLERECVLEWLRYLEMGKETT 203
QY 181 DQOVPLPVKVVTHVTS-VTTLCRALNYYPNITMKWLKDKQPMDAKEPFEPKQVLPNGD 239
DB 204 QRADPPKAHVTHHPASDREATLRCWALGFYPAISLTWQDGED-QTQDTLVELVTRPGD 262
QY 240 GTYQGHMITLAVPGEQRQYTCQVEHPGLDQPLVIWE 276
DB 263 GTFQKAAVVPVSGEQRQYTCRVOHEGLPEPLTLTWE 299

RESULT 4
I46858
MHC class I RLA precursor - rabbit
C:Species: Oryctolagus cuniculus (domestic rabbit)
C:Date: 14-Feb-1997 #sequence_revision 14-Feb-1997 #text_change 21-Jan-2000
C:Accession: I46858
R:Marche, P.N.; Tykocinski, M.L.; Max, E.E.; Kindt, T.J.
Immunogenetics 21, 71-82, 1985
A:Title: Structure of a functional rabbit class I MHC gene: Similarity to human
A:Reference number: I46858; MUID:85103547; PMID:3917974
A:Accession: I46858
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-361 <VAR>
A:Cross-references: GB:K02819; NID:g165497; PIDN:AAA98730.1; PID:g165498
C:Genetics:
A:Introns: 25/1; 115/1; 207/1; 295/1; 337/1; 348/1
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 34.1%; Score 517; DB 2; Length 361;
Best Local Similarity 40.1%; Pred. No. 1.7e-35;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7

QY 5 SHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDYHE--SRRVEPTPWSSRISSQW 62
DB SHSRYFYTSVRPGLGEPRFIIVGYDDTQVRFDSDAASPRMEQAPWM-GQVEPEYW 84
QY 63 LQLSOSLKGWDHMTVDFTWIMENHNASKB-SHTLQVILGCEMQEDNS-TGYWKYGYDG 120
DB DQQTQIAKTAQTFRVNLNTALRYNQSAAGSHTFQTMFGCEVWADGRFFHGYQYAYDG 144
QY 121 QDALEFCPDTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQLLELGRGVL 180
DB 145 ADVIALNEDLRSWTAADTAQNTQKWEAAG-EAERHRAVLERECVLEWLRYLEMGKETT 203
QY 181 DQOVPLPVKVVTHVTS-VTTLCRALNYYPNITMKWLKDKQPMDAKEPFEPKQVLPNGD 239
DB 204 QRADPPKAHVTHHPASDREATLRCWALGFYPAISLTWQDGED-QTQDTLVELVTRPGD 262
QY 240 GTYQGHMITLAVPGEQRQYTCQVEHPGLDQPLVIWE 276
DB 263 GTFQKAAVVPVSGEQRQYTCRVOHEGLPEPLTLTWE 299

RESULT 5

S06424
MHC class I histocompatibility antigen Ch25 alpha chain precursor - chimpanzee
N;Alternate names: MHC Ch1A chain
C;Species: Pan troglodytes (chimpanzee)
C;Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 23-Jul-1999
C;Accession: S06424; I36959
R;Lawlor, D.A.; Ward, F.E.; Ennis, P.D.; Jackson, A.P.; Parham, P.
Nature 335, 268-271, 1988
A;Title: HLA-A and B polymorphisms predate the divergence of humans and chimpanzees.
A;Reference number: S06424; MUID:98319000; PMID:3412487
A;Accession: S06424
A;Molecule type: mRNA
A;Residues: 1-332 <LAW>
R;Parham, P.; Lawlor, D.A.; Lomen, C.E.; Ennis, P.D.
J. Immunol. 142, 3937-3950, 1989
A;Title: Diversity and diversification of HLA-A,B,C alleles.
A;Reference number: I36956; MUID:89235215; PMID:2715640
A;Accession: I36959
A;Molecule type: mRNA
A;Residues: 1-332 <RES>
A;Cross-references: GB:M24047; NID:g176818; PID:AAA35426.1; PID:g553155
C;Superfamily: Class I histocompatibility antigen; immunoglobulin homology
C;Keywords: glycoprotein; membrane protein
F;1-24/Domain: signal sequence #status predicted <SIG>
F;25-114/Domain: alpha-1 #status predicted <EX1>
F;115-206/Domain: alpha-2 #status predicted <EX2>
F;220-285/Domain: immunoglobulin homology <IMM>
F;307-331/Domain: transmembrane #status predicted <TM>
F;110/Binding site: carbohydrate (Asn) (covalent) #status predicted
F;125-188, 227-283/Disulfide bonds: #status predicted

Query Match 33.9%; Score 514; DB 2; Length 332;
Best Local Similarity 40.1%; Pred. No. 2.7e-35;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;

QY 5 SLSHLVFMGASEQDGLSLFEALGVYDDQLFVFDHE--SRVRPRTPMVSSRISSQMW 62

DB 26 SHSMRYFFTSVSRPGRGEPRFIAVGVDYDTQFVFDSDAASQRMPEAPWIEQ-GPEY 84

QY 63 LQLSQLKGDHMFVTVDFTWIMENHNASKE-SHTLQVILGCEMOEDNS-TEGYWKYGYDG 120

DB 85 DQTRAKAHSQTRDVLGLTRGYNQSDGSHTIQIMYGCDDVGSGRFLRGTRQDAYDG 144

QY 121 QDALEFCPTDLWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

DB 145 KDVIALLNEDLRSWTAADMAAQITKRKEAAH-AAEQRAYLEGTVCVWELRRYLENGKEYL 203

QY 181 DQOVPLVKTTH-VTSSVTLRCALNYYPQNIWKWKLDKQPMDAKFEPEKDVLPNGD 239

DB 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLVELVETRPAGD 262

QY 240 GTYQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 276

DB 263 GTFOKAAVVPVSGEQRVYCHVQHEGLPKPLTLRWE 299

RESULT 6

I36961

MHC class I protein - chimpanzee

C;Species: Pan troglodytes (chimpanzee)

C;Date: 04-Oct-1996 #sequence_revision 04-Oct-1996 #text_change 21-Jan-2000

C;Accession: I36961

R;Lawlor, D.A.; Ward, F.E.; Ennis, P.D.; Jackson, A.P.; Parham, P.

Immunol. Rev. 113, 147-185, 1990

A;Title: Comparison of class I MHC alleles in humans and apes.

A;Reference number: I36961; MUID:90201944; PMID:1690682

A;Accession: I36961

A;Status: preliminary; translated from GB/EMBL/DDBJ

A;Molecule type: mRNA

A;Residues: 1-365 <RES>

A;Cross-references: GB:M30678; NID:g176822; PID:AAA87970.1; PID:g176823

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 33.9%; Score 514; DB 2; Length 332;
Best Local Similarity 40.1%; Pred. No. 2.7e-35;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;

QY 5 SLSHLVFMGASEQDGLSLFEALGVYDDQLFVFDHE--SRVRPRTPMVSSRISSQMW 62

DB 26 SHSMRYFFTSVSRPGRGEPRFIAVGVDYDTQFVFDSDAASQRMPEAPWIEQ-GPEY 84

QY 63 LQLSQLKGDHMFVTVDFTWIMENHNASKE-SHTLQVILGCEMOEDNS-TEGYWKYGYDG 120

DB 85 DQTRAKAHSQTRDVLGLTRGYNQSDGSHTIQIMYGCDDVGSGRFLRGTRQDAYDG 144

QY 121 QDALEFCPTDLWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

DB 145 KDVIALLNEDLRSWTAADMAAQITKRKEAAH-AAEQRAYLEGTVCVWELRRYLENGKEYL 203

QY 181 DQOVPLVKTTH-VTSSVTLRCALNYYPQNIWKWKLDKQPMDAKFEPEKDVLPNGD 239

DB 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLVELVETRPAGD 262

QY 240 GTYQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 276

DB 263 GTFOKAAVVPVSGEQRVYCHVQHEGLPKPLTLRWE 299

RESULT 7

B27638

MHC class I histocompatibility antigen alpha chain precursor (BL3-7) - bovine

C;Species: Bos primigenius taurus (cattle)

C;Date: 08-Mar-1989 #sequence_revision 08-Mar-1989 #text_change 16-Feb-1997

C;Accession: B27638

R;Ennis, P.D.; Jackson, A.P.; Parham, P.

J. Immunol. 141, 642-651, 1988

A;Title: Molecular cloning of bovine class I MHC cDNA.

A;Reference number: A92826; MUID:89258075; PMID:3133413

A;Accession: B27638

A;Status: not compared with conceptual translation

A;Molecule type: mRNA

A;Residues: 1-361 <ENN>

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

C;Keywords: heterodimer; transmembrane protein

F;1-24/Domain: signal sequence #status predicted <SIG>

F;25-361/Product: MHC class I histocompatibility antigen, BoLA alpha chain (BL3-7) #stat

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.7%; Score 510; DB 2; Length 361;
Best Local Similarity 38.9%; Pred. No. 6.4e-35;
Matches 109; Conservative 49; Mismatches 114; Indels 8; Gaps 7;

QY 2 LRLSHLVFMGASEQDGLSLFEALGVYDDQLFVFDHE--SRVRPRTPMVSSRISS 59

DB 23 LAGSHLVYFTYGVSRPGLGEPRFIAVGVDYDTQFVFDSDAPNPREPVRVMEQEQ-GP 81

QY 60 QMWLQSLKGDHMFVTVDFTWIMENHNASKE-SHTLQVILGCEMOEDNS-TEGYWKY 117

DB 82 EYWDNTRIYKDTAQIFRVDLNLTRGYNQSGTSHNQAMYGCDVGDGPDGRLLRGFWQFG 141

QY 118 YDQDALEFCPTDLWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGR 177

DB 142 YDGRDYIALNEELRSWTAADTAQAQITKRKEAAH-AAETWRNLYEGECVWELRRYLENGK 200

QY 178 GVLDDQVPLVKTTH-VTSSVTLRCALNYYPQNIWKWKLDKQPMDAKFEPEKDVLP 236

DB 201 DTLRLADPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDMELVETRP 259

QY 237 NGDGYTQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 276

DB 260 SDGTGTFQKAAVVPVSGEQRVYCHVQHEGLPKPLTLRWE 299

RESULT 8

A>Note: this allele is designated A*1101 (formerly AllE, All.1)
R;Lin, L.; Tokunaga, K.; Ishikawa, Y.; Bannai, M.; Kashiwase, K.; Kuwata, S.; Akaza, T.
Tissue Antigens 43, 78-82, 1994
A>Title: Sequence analysis of serological HLA-A11 split antigens, All.1 and All.2.
A;Reference number: I60129; MUID:94287401; PMID:16016945
A;Accession: I60129
A;Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A;Residues: 1-365 <RES>
A;Cross-references: GB:M16841; NID:G540516; PIDN:BAA04117.1; PID:9487909
A>Note: this allele is designated A*1101 (formerly AllE, All.1)
C;Genetics:
A;Gene: GDB:HLA-A
A;Cross-references: GDB:119310; OMIM:142800
A;Map position: 6p21.3-6p21.3
C;Superfamily: class I histocompatibility antigen; immunoglobulin homology
C;Keywords: transmembrane protein
F;1-24/Domain: signal sequence #status predicted <SIG>
F;25-365/Product: class I histocompatibility antigen alpha chain #status predicted <MAT>
F;25-298/Domain: extracellular #status predicted <EXT>
F;220-285/Domain: immunoglobulin homology <IMM>
F;299-337/Domain: transmembrane #status predicted <TMW>
F;338-365/Domain: intracellular #status predicted <INT>

Query Match 33.6%; Score 508; DB 2; Length 365;
Best Local Similarity 39.4%; Pred.No.9.5e-35;
Matches 109; Conservative 46; Mismatches 114; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGVVDQLFVFDHE--SRVEPRTPWVSRRISQQW 62
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :
Db 26 SHSMRYFYTSVRPGRGEPFRFTAVGVVDPTQVFPSDASQRMEPAPWBOE-GPEYW 84
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :
QY 63 LOLSQSLLKGDHMFVTDFWTIENHNASKS-SHTLQVLGCMEQDNS-TGYWKYGYDG 120
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :
Db 85 DQETRNVAQSQTDRVDLTGLTGAGYNQSDGSHTTQIMYGCDVGDPGRFLGYRDAYDG 144
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :
QY 121 QDALEFCPTLDWRAPRAWPPTYKLEWERHKIRARQNRAYLERDCPAQIQQLLELRGVL 180
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :
Db 145 KDVIALLNEDLRSWTAADMAAQITKRWEAAH-AAEQRAYLEGRCVLEWRLRYLENGKETL 203
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :
QY 181 DQGVPLVKVTHH-VTSSVTTLRCRALNYYPONIIMKLKDQPMDAKEFPKDVLPNGD 239
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :
Db 204 QRTDDPKTMTHTPHSIDSEATLRCWALGFYPAEITLTWORDGED-QTQDELVETRPA GD 262
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :
QY 240 GTVQGMITLAVPPGEORVTCOVERHPGLDPLIVIVE 276
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :
Db 263 GTFGKAAVVPSGEGEORTCHVQHEGLPKPIILRWE 299
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :

RESULT 10
I56039
HLA-A30.3 precursor - human
C;Species: Homo sapiens (man)
C;Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 21-Jan-2000
C;Accession: I56039
R;Kato, K.; Trapana, J.A.; Allopenna, J.; Dupont, B.; Yang, S.Y.
J. Immunol. 143, 3371-3378, 1989
A>Title: Molecular analysis of the serologically defined HLA-Aw19 antigens. A genetical
A;Reference number: I56039; MUID:90038496; PMID:2478623
A;Accession: I56039
A;Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A;Residues: 1-365 <RES>
A;Cross-references: GB:M30576; NID:g187646; PIDN:AAA59612.1; PID:9386978
C;Superfamily: class I histocompatibility antigen; immunoglobulin homology
F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.6%; Score 508; DB 2; Length 365;
Best Local Similarity 39.4%; Pred.No.9.5e-35;
Matches 109; Conservative 47; Mismatches 113; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGVVDQLFVFDHE--SRVEPRTPWVSRRISQQW 62
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :
||| : | : | : | : | : | : | : | : | : | : | : | : | : | :

```
Db 26 SHSMRYFFTSVRPGSGEPRIAGVYDDTQFVRFSDSDAASQRMPEPRAPWIEQ-EPEYW 84
QY 63 LQLSLSKLGWDHMTFVDFTWIMENHNASK-SHTLQVILGCEMQEDNS--TEGYWKYGYDG 120
Db 85 DQETRNVAQSQDTRVDLGLTGRYVNSQSEAGSHTIQIMYGCDVSGDRFLRGVYEQHAYDG 144
QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHKKIRARQNRAYLERDCPAQLQQLLELGRGV 180
Db 145 KDYIALNEDLRSWTAADMAAQITQRKWEAAR-WAEQLRAYLEGTCVWELRYLENGKETL 203
QY 181 DQOVPPPLVKVTHH-VTSSVTLRCALNYYPONITMKWLKDKQPMDAKEFEKPDVLPNGD 239
Db 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQRDGED-QTQDTLVELTRPADG 262
QY 240 GTYQGWITLAVPPGEQRYTCQVEHPGLDQPLIVWE 276
Db 263 GTFQKAAVAVVPSGGEQRYTCHVQHEGLPKPLTLRWE 299

RESULT 11
HLHUA3
MHC class I histocompatibility antigen HLA-A3 alpha chain precursor - human
C:Species: Homo sapiens (man)
C>Date: 17-Mar-1987 #sequence_revision 17-Mar-1987 #text_change 02-Sep-1997
R:Accession: A02192
R:Strachan, T.; Sodoyer, R.; Damotte, M.; Jordan, B.R.
EMBO J. 3, 887-894, 1984
A:Title: Complete nucleotide sequence of a functional class I HLA gene, HLA-A3: implicated
A:Reference number: A02192; MUID:84207948; PMID:6609814
A:Accession: A02192
A:Molecule type: DNA
A:Residues: 1-370 <STR>
C:Genetics:
A:Gene: GDB:HLA-A
A:Cross-references: GDB:119310; OMIM:142800
A:Map position: 6p21.3-6p21.3
A:Introns: 30/1; 120/1; 212/1; 304/1; 343/1; 354/1; 370/1
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
C:Keywords: duplication; glycoprotein; heterodimer; transmembrane protein; transplantati
F;1-29/Domain: signal sequence #status predicted <SIG>
F;30-370/Product: class I histocompatibility antigen HLA-A3 alpha chain #status predicte
F;30-312/Domain: extracellular #status predicted <EX1>
F;30-119/Domain: alpha-1 <EX2>
F;120-211/Domain: alpha-2 <EX2>
F;225-290/Domain: immunoglobulin homology <IMM>
F;313-337/Domain: transmembrane #status predicted <TM>
F;338-370/Domain: intracellular #status predicted <INT>
F;115/Binding site: carbohydrate (Asn) (covalent) #status predicted
F;232-288/Disulfide bonds: #status predicted

Query Match 33.4%; Score 506; DB 1; Length 370;
Best Local Similarity 39.6%; Pred. No. 1.4e-34;
Matches 110; Conservative 46; Mismatches 112; Indels 10; Gaps 8;

QY 5 SHSLHYLFMGASQDGLSLFEALGVYDDQLFVFDHE--SRVPRTPWSSRISSQW 62
Db 31 SHSMRYFFTSVRPGSGEPRIAGVYDDTQFVRFSDSDAASQRMPEPRAPWIEQ-EPEYW 89

QY 63 LQLSLSKLGWDHMTFVDFTWIMENHNASK-SHTLQVILGCEMQEDNS--TEGYWKYGYDG 120
Db 90 DQETRNVAQSQDTRVDLGLTGRYVNSQSEAGSHTIQIMYGCDVSGDRFLRGVYEQHAYDG 149
QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHKKIRARQNRAYLERDCPAQLQQLLELGRGV 179
Db 150 KDYIALNEDLRSWTAADMAAQITQRKWEAAR-WAEQLRAYLEGTCVWELRYLENGKET 207
QY 180 LDQOVPPPLVKVTHH-VTSSVTLRCALNYYPONITMKWLKDKQPMDAKEFEKPDVLPNG 238
Db 208 LQRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQRDGED-QTQDTLVELTRPADG 266
QY 239 DGTQGWITLAVPPGEQRYTCQVEHPGLDQPLIVWE 276
Db 267 DGTQKAAVAVVPSGGEQRYTCHVQHEGLPKPLTLRWE 304
```

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RESULT 12
I38439
MHC class I histocompatibility antigen HLA-A*8001 precursor - human
C:Species: Homo sapiens (man)
C>Date: 07-Jun-1996 #sequence_revision 07-Jun-1996 #text_change 21-Jan-2000
C:Accession: I59638; I38439
R:Domena, J.D.; Hildebrand, W.H.; Bias, W.B.; Parham, P.
Tissue Antigens 42, 156-159, 1993
A:Title: A sixth family of HLA-A alleles defined by HLA-A*8001.
A:Reference number: I59638; MUID:94112691; PMID:8284791
A:Accession: I59638
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-365 <DOM>
A:Cross-references: GB:118898; NID:G306853; PIDN:AAAL7012.1; PID:G306854
R:Balas, A.; Garcia-Sanchez, F.; Gomez-Reino, F.; Vicario, J.L.
Immunogenetics 39, 452, 1994
A:Title: Characterization of a new and highly distinguishable HLA-A allele in a Spanish
A:Reference number: I38439; MUID:94245293; PMID:8188325
A:Accession: I38439
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-365 <BAL>
A:Cross-references: EMBL:U03754; NID:G432407; PIDN:AAC04322.1; PID:G432408
C:Genetics:
A:Gene: GDB:HLA-A
A:Cross-references: GDB:119310; OMIM:142800
A:Map position: 6p21.3-6p21.3
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.3%; Score 504; DB 2; Length 365;
Best Local Similarity 38.3%; Pred. No. 2.1e-34;
Matches 106; Conservative 52; Mismatches 111; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDGLSLFEALGVYDDQLFVFDHE--SRVPRTPWSSRISSQW 62
Db 26 SHSMRYFFTSVRPGSGEPRIAGVYDDTQFVRFSDSDAASQRMPEPRAPWIEQ-EPEYW 84

QY 63 LQLSLSKLGWDHMTFVDFTWIMENHNASK-SHTLQVILGCEMQEDNS--TEGYWKYGYDG 120
Db 85 DEETRNVAQSQDTRVDLGLTGRYVNSQSEAGSHTIQIMYGCDVSGDRFLRGVYEQHAYDG 144
QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHKKIRARQNRAYLERDCPAQLQQLLELGRGV 180
Db 145 KDYIALNEDLRSWTAADMAAQITQRKWEAAR-WAEQLRAYLEGTCVWELRYLENGKETL 203
QY 181 DQOVPPPLVKVTHH-VTSSVTLRCALNYYPONITMKWLKDKQPMDAKEFEKPDVLPNGD 239
Db 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQRDGED-QTQDTLVELTRPADG 262
QY 240 GTYQGWITLAVPPGEQRYTCQVEHPGLDQPLIVWE 276
Db 263 GTFQKAAVAVVPSGGEQRYTCHVQHEGLPKPLTLRWE 299

RESULT 13
I37542
MHC class I histocompatibility antigen HLA-A2 alpha chain (allele A*0216) precursor - h
C:Species: Homo sapiens (man)
C>Date: 04-Oct-1996 #sequence_revision 04-Oct-1996 #text_change 21-Jan-2000
C:Accession: I37542; 849582
R:Barouch, D.; Krausa, P.; Bodmer, J.; Browning, M.J.; McMichael, A.J.
Immunogenetics 41, 388, 1995
A:Title: Identification of a novel HLA-A2 subtype, HLA-A*0216.
A:Reference number: I37542; MUID:95278976; PMID:7759139
A:Accession: I37542
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-365 <RES>
A:Cross-references: EMBL:Z46633; NID:G575248; PIDN:CAA86602.1; PID:G575249
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A;Note: submitted to the EMBL Data Library, November 1994

C;Genetics:

A;Gene: hla-A

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.2%; Score 503; DB 2; Length 365;
Best Local Similarity 39.4%; Pred. No. 2.5e-34;
Matches 109; Conservative 45; Mismatches 115; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDGLSLFEALGVYDDQLFVFDHE--SRVPRTPWSSRISSQMW 62

Db 26 SHSMRYFTTSVRGEPGFIAVGVDQTFVRFDSDAASQRMPEAPWIEQ-GPEYW 84

QY 63 LQLSLSLKGWDHMTFTVDFTIMENHNASKB-SHTLQVILGCEMQED-NSTEGYWKYGYDG 120

Db 85 DGETRKVKVKAHSQTHRVDLGLTRGYNQSEAGSHTVQRMVGCVDGSDWRFLRGVHQYAYDG 144

QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180

Db 145 KDYIALKEDLRSWTAADMAAQTTHKWEAAHV-AEQRLAYLEGCEVWLRRLRYLENGKETL 203

QY 181 DQOVPLVKVTHH-VTSSVTLRCRALNYPQNTIMKWLKQKQMDAKFEKPKDVLNPGD 239

Db 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPABEITLTWQDGDG-QTQDTLVETRPAGD 262

QY 240 GTYQGWITLAVPGEQRVTCVHEHPCLDQPLIVWE 276

Db 263 GTFOKAAVVPVSGQEQRYTCHVQHEGLPKPLTLRWE 299

RESULT 14

I38442

Gene HLA-A-0205 protein - human

C;Species: Homo sapiens (man)

C;Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 21-Jan-2000

C;Accession: I38442

R;Holmes, N.; Ennis, P.; Wan, A.M.; Denney, D.W.; Parham, P.

J. Immunol. 139, 936-941, 1987

A;Title: Multiple genetic mechanisms have contributed to the generation of the HLA-A2/A2

A;Reference number: I38441; MUID:87252273; PMID:3496393

A;Status: preliminary; translated from GB/EMBL/DDBJ

A;Molecule type: DNA

A;Residues: 1-365 <RES>

A;Cross-references: EMBL:U03862; NID:9432436; PIDN:AAA03603.1; PID:9432437

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.2%; Score 503; DB 2; Length 365;
Best Local Similarity 39.7%; Pred. No. 2.5e-34;
Matches 110; Conservative 43; Mismatches 116; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDGLSLFEALGVYDDQLFVFDHE--SRVPRTPWSSRISSQMW 62

Db 26 SHSMRYFTTSVRGEPGFIAVGVDQTFVRFDSDAASQRMPEAPWIEQ-GPEYW 84

QY 63 LQLSLSLKGWDHMTFTVDFTIMENHNASKB-SHTLQVILGCEMQED-NSTEGYWKYGYDG 120

Db 85 DGETRKVKVKAHSQTHRVDLGLTRGYNQSEAGSHTVQRMVGCVDGSDWRFLRGVHQYAYDG 144

QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180

Db 145 KDYIALKEDLRSWTAADMAAQTTHKWEAAHV-AEQWRAYLEGTCVWLRRLRYLENGKETL 203

QY 181 DQOVPLVKVTHH-VTSSVTLRCRALNYPQNTIMKWLKQKQMDAKFEKPKDVLNPGD 239

Db 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPABEITLTWQDGDG-QTQDTLVETRPAGD 262

QY 240 GTYQGWITLAVPGEQRVTCVHEHPCLDQPLIVWE 276

Db 263 GTFOKAAVVPVSGQEQRYTCHVQHEGLPKPLTLRWE 299

RESULT 15

I61902

MHC class I histocompatibility antigen HLA-A alpha chain precursor - human (isolate A*0.

C;Species: Homo sapiens (man)

A;Variety: isolate A*0212

C;Date: 06-Sep-1996 #sequence_revision 06-Sep-1996 #text_change 23-Jul-1999

C;Accession: I61902

R;Belich, M.P.; Madrigal, J.A.; Hildebrand, W.H.; Zemmour, J.; Williams, R.C.; Luz, R.;

Nature 357, 326-329, 1992

A;Title: Unusual HLA-B alleles in two tribes of Brazilian Indians.

A;Reference number: I37120; MUID:92269955; PMID:1317015

A;Accession: I61902

A;Status: translated from GB/EMBL/DDBJ

A;Molecule type: mRNA

A;Residues: 1-365 <RES>

A;Cross-references: GB:M84378; NID:9187625; PIDN:AAA59604.1; PID:9187626

A;Experimental source: cell line KRC 033; isolate A*0212

C;Genetics:

A;Gene: GDB:HLA-A

A;Cross-references: GDB:119310; OMIM:142800

A;Map position: 6p21.3-6p21.3

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;1-24/Domain: signal sequence protein

F;25-365/Product: MHC class I histocompatibility antigen HLA-A alpha chain #status pred.

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.2%; Score 503; DB 2; Length 365;
Best Local Similarity 39.4%; Pred. No. 2.5e-34;
Matches 109; Conservative 44; Mismatches 116; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDGLSLFEALGVYDDQLFVFDHE--SRVPRTPWSSRISSQMW 62

Db 26 SHSMRYFTTSVRGEPGFIAVGVDQTFVRFDSDAASQRMPEAPWIEQ-GPEYW 84

QY 63 LQLSLSLKGWDHMTFTVDFTIMENHNASKB-SHTLQVILGCEMQED-NSTEGYWKYGYDG 120

Db 85 DGETRKVKVKAHSQTHRVDLGLTRGYNQSEAGSHTVQRMVGCVDGSDWRFLRGVHQYAYDG 144

QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180

Db 145 KDYIALKEDLRSWTAADMAAQTTHKWEAAHV-AEQRAYLEGTCVWLRRLRYLENGKETL 203

QY 181 DQOVPLVKVTHH-VTSSVTLRCRALNYPQNTIMKWLKQKQMDAKFEKPKDVLNPGD 239

Db 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPABEITLTWQDGDG-QTQDTLVETRPAGD 262

QY 240 GTYQGWITLAVPGEQRVTCVHEHPCLDQPLIVWE 276

Db 263 GTFOKAAVVPVSGQEQRYTCHVQHEGLPKPLTLRWE 299

Search completed: August 5, 2003, 13:10:37

Job time : 16 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:05:29 ; Search time 9.5 Seconds
(without alignments)
1366.250 Million cell updates/sec

Title: US-10-092-404-3

Perfect score: 1514

Sequence: 1 RLRLSHLHFLWFGASEQDL.....RYTCQVEHFGLDQPLIVWE 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SwissProt_41.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1502	99.2	348	1	HFE HUMAN
2	1227	81.0	348	1	HFE_DICSU
3	1225	80.9	348	1	HFE_CERSI
4	1221	80.6	348	1	HFE_RHIUN
5	1218	80.4	348	1	HFE_DICBI
6	1145	75.6	360	1	HFE_RAT
7	1129	74.6	359	1	HFE_MOUSE
8	517	34.1	361	1	HALA_RABIT
9	517	34.1	361	1	HALB_RABIT
10	511	33.8	365	1	LA01_PANTR
11	510	33.7	364	1	HALB_BOVIN
12	508	33.6	365	1	LA01_HUMAN
13	506	33.4	370	1	LA03_HUMAN
14	504	33.3	365	1	LA80_HUMAN
15	502	33.2	365	1	LA30_HUMAN
16	500	33.0	365	1	LA02_HUMAN
17	500	33.0	365	1	LA30_HUMAN
18	500	33.0	365	1	LA03_HUMAN
19	498	32.9	365	1	LA03_PANTR
20	497	32.8	365	1	LA33_HUMAN
21	497	32.8	365	1	LA36_HUMAN
22	497	32.8	365	1	LA68_HUMAN
23	495.5	32.7	362	1	HAL9_CANFA
24	495	32.7	365	1	LA01_HUMAN
25	494	32.6	273	1	LA69_HUMAN
26	494	32.6	296	1	ZA2G_RAT
27	494	32.6	365	1	LA04_PANTR
28	494	32.6	365	1	LA24_HUMAN
29	492	32.5	360	1	HALA_BOVIN
30	491	32.4	362	1	LB45_HUMAN
31	490	32.4	365	1	LA23_HUMAN
32	488	32.2	338	1	LB20_HUMAN
33	487	32.2	363	1	LB04_GORGO

34	486	32.1	295	1	ZA2G_HUMAN	P25311 homo sapien
35	486	32.1	322	1	HA10_MOUSE	P01898 mus musculus
36	486	32.1	362	1	IB29_HUMAN	P18463 homo sapien
37	486	32.1	371	1	HA12_RAT	P16391 rattus norv
38	485	32.0	365	1	LA34_HUMAN	P30453 homo sapien
39	485	32.0	365	1	LA66_HUMAN	P30457 homo sapien
40	484	32.0	338	1	HLAG_HUMAN	P17693 homo sapien
41	484	32.0	361	1	IB14_HUMAN	P03989 homo sapien
42	484	32.0	362	1	IB18_HUMAN	P10318 homo sapien
43	484	32.0	366	1	IC02_GORGO	P30385 gorilla gor
44	484	32.0	366	1	IC04_GORGO	P30387 gorilla gor
45	483	31.9	359	1	IB01_PANTR	P13750 pan troglod

ALIGNMENTS

RESULT 1
HFE_HUMAN
ID HFE_HUMAN STANDARD; PRT; 348 AA.
AC Q30201; O75929; O75930; Q96KU5; Q96KU7; Q96KU8; Q96KU9; Q96KU6;
AC Q9HC68; Q9HC70; Q9HC83;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Hereditary hemochromatosis protein precursor (HLA-H).
GN HFE OR HLAH.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RX MEDLINE=963311279; PubMed=8696333;
RA Feder J.N., Gnirke A., Thomas W., Tsuchihashi Z., Ruddy D.A.,
RA Hinton L.M., Jones N.L., Kimmel B.E., Kronmal G.S., Lauer P.,
RA Basava A., Dornishian F., Domingo R., Ellis M.C. Jr., Fullan A.,
RA Lee V.K., Loeb D.B., Mapa F.A., McClelland E., Meyer N.C.,
RA Mintier G.A., Moeller N., Moore T., Morikang E., Prass C.E.,
RA Quintana L., Starnes S.M., Schatzman R.C., Brunke K.J.,
RA Drayna D.T., Risch N.J., Bacon B.R., Wolff R.K.;
RT "A novel MHC class I-like gene is mutated in patients with hereditary
haemochromatosis.";
RL Nat. Genet. 13:399-409(1996).
SEQUENCE FROM N.A. (ISOFORM 1).
ALBIG W., Burmester N., Bode C., Doenecke D., Drabent B.;
Submitted (MAR-1997) to the EMBL/GenBank/DBJ databases.
SEQUENCE FROM N.A. (ISOFORM 1).
Ruddy D.A., Kronmal G.S., Lee V.K., Mintier G.A., Quintana L.,
Domingo R. Jr., Meyer N.C., Irrinki A., McClelland E.E., Fullan A.,
Mapa F.A., Moore T., Thomas W., Loeb D.B., Harmon C., Tsuchihashi Z.,
Wolff R.K., Schatzman R.C., Feder J.N.;
RT "A 1.1-Mb transcript map of the hereditary hemochromatosis locus.";
RL Genome Res. 7:441-456(1997).
SEQUENCE FROM N.A. (ISOFORM 1).
Gasparini P.,
Submitted (SEP-1997) to the EMBL/GenBank/DBJ databases.
SEQUENCE FROM N.A. (ISOFORMS 2; 3 AND 4).
MEDLINE=99180629; PubMed=10079302;
RA Rhodes D.A., Towdale J.;
RT "Alternate splice variants of the hemochromatosis gene Hfe.";
RL Immunogenetics 49:357-359(1999).
SEQUENCE FROM N.A. (ISOFORMS 2; 5; 6 AND 7).
Oliva R., Sanchez M.;
RT "Identification of different alternative splicing forms of the HFE
gene.";
Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.

[7] SEQUENCE FROM N.A. (ISOFORMS 1; 7; 8; 9 AND 10).
RP MEDLINE=20448010; PubMed=11001625;
RX Thénie A., Orhant M., Gicquel I., Fergelot P., Le Gall J.-Y.,
RA David V., Mosser J.;
RT "The HFE gene undergoes alternate splicing processes.";
RL Blood Cells Mol. Dis. 26:155-162(2000).
RN [8]
RP FUNCTION.
RX MEDLINE=98132614; PubMed=9465039;
RA Feder J.N., Penny D.M., Irrinki A., Lee V.K., Lebron J.A., Watson N.,
RX Tsuchihashi Z., Sigal E., Bjorkman P.J., Schatzman R.C.;
RA "The hemochromatosis gene product complexes with the transferrin
RT receptor and lowers its affinity for ligand binding.";
RL Proc. Natl. Acad. Sci. U.S.A. 95:1472-1477(1998).
RN [9]
RP X-RAY CRYSTALLOGRAPHY (2.6 ANGSTROMS).
RX MEDLINE=98206473; PubMed=9546397;
RA Lebron J.A., Bennett M.J., Vaughn D.E., Chirino A.J., Snow P.M.,
RX Mintier G.A., Feder J.N., Bjorkman P.J.;
RA "Crystal structure of the hemochromatosis protein HFE and
RT characterization of its interaction with transferrin receptor.";
RL Cell 93:111-123(1998).
RN [10]
RP VARIANTS HH ASP-63 AND TYR-282.
RX MEDLINE=97260408; PubMed=9106528;
RA Carella M., D'Ambrosio L., Totaro A., Grifa A., Valentino M.A.,
RA Piperno A., Girelli D., Roetto A., Franco B., Gasparini P.,
RA Camaschella C.;
RT "Mutation analysis of the HLA-H gene in Italian hemochromatosis
patients.";
RL Am. J. Hum. Genet. 60:828-832(1997).
RN [11]
RP VARIANT HH/PCT TYR-282.
RX MEDLINE=97176837; PubMed=9024376;
RA Roberts A.G., Whitley S.D., Morgan R.R., Worwood M., Elder G.H.;
RT "Increased frequency of the hemochromatosis Cys282Tyr mutation in
RL sporadic porphyria cutanea tarda.";
RN Lancet 349:321-323(1997).
RX MEDLINE=98085904; PubMed=9425935;
RA Sampietro M., Piperno A., Lupica L., Arosio C., Vergani A.,
RA Corbetta N., Malosio I., Mattioli M., Fracanzani A.L.,
RA Cappellini M.D., Fiorelli G., Fargion S.;
RT "High prevalence of the Hs63Asp HFE mutation in Italian patients with
RL porphyria cutanea tarda.";
RN Hepatology 27:181-184(1998).
RX VARIANTS HH/PCT ASP-63 AND TYR-282.
RX MEDLINE=98281650; PubMed=9620340;
RA Bonkovsky H.L., Poh-Fitzpatrick M., Pinstone N., Obando J.,
RA Di Bisceglie A., Tattire C., Tortorelli K., LeClair P., Mercurio M.G.,
RA Lambrecht R.W.;
RT "Porphyria cutanea tarda, hepatitis C, and HFE gene mutations in North
RL America.";
RN Hepatology 27:1661-1669(1998).
RX VARIANTS HH ASP-63; CYS-65 AND TYR-282.
RX MEDLINE=99211934; PubMed=10194428;
RA Mura C., Ragues O., Ferec C.;
RT "HFE mutations analysis in 711 hemochromatosis probands: evidence for
RL S65C implication in mild form of hemochromatosis.";
RN Blood 93:2502-2505(1999).
RX VARIANTS HH CYS-65; ARG-93 AND THR-105.
RX MEDLINE=20042794; PubMed=10575540;
RA Barton J.C., Sawada-Hirai R., Rothenberg B.E., Acton R.T.;
RT "Two novel missense mutations of the HFE gene (I105T and G93R) and
RL identification of the S65C mutation in Alabama hemochromatosis
proband.";
RN Blood Cells Mol. Dis. 25:147-155(1999).
RN

RP VARIANTS VP ASP-63 AND HIS-127, VARIANT HH MET-330, AND VARIANTS
RP MET-53 AND MET-59.
RX MEDLINE=99330560; PubMed=10401000;
RA de Villiers J.N.P., Hillermann R., Loubser L., Kotze M.J.;
RT "Spectrum of mutations in the HFE gene implicated in haemochromatosis
and porphyria.";
RL Hum. Mol. Genet. 8:1517-1522(1999).
RN [17]
RP VARIANTS HH ASP-63 AND TYR-282.
RX MEDLINE=99140360; PubMed=10094552;
RA Merryweather-Clarke A.T., Simonsen H., Shearman J.D., Pointon J.J.,
RA Norgaard-Pedersen B., Robson K.J.H.;
RT "A retrospective anonymous pilot study in screening newborns for HFE
RL mutations in Scandinavian populations.";
RN Hum. Mutat. 13:154-159(1999).
RN [18]
RP VARIANT HH CYS-65.
RX Fagan E., Payne S.J.;
RA "A novel missense mutation S65C in the HFE gene with a possible role
RT in hereditary haemochromatosis.";
RL Hum. Mutat. 13:507-508(1999).
RN [19]
RP VARIANT LYS-277.
RX MEDLINE=20081073; PubMed=10612845;
RA Bradbury R., Fagan E., Payne S.J.;
RT "Two novel polymorphisms (E277K and V212V) in the haemochromatosis
RL gene HFE.";
RN Hum. Mutat. 15:120-120(2000).
CC -!- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
CC affinity for iron-loaded transferrin.
CC -!- SUBCELLULAR LOCATION: Type I membrane protein.
CC -!- ALTERNATIVE PRODUCTS:
CC Event=Alternative splicing; Named isoforms=10;
CC Comment=Additional isoforms seem to exist;
CC Name=1;
CC IsoId=Q30201-1; Sequence=Displayed;
CC Name=2; Synonyms=delE2;
CC IsoId=Q30201-2; Sequence=VSP_003218;
CC Name=3; Synonyms=del14B4;
CC IsoId=Q30201-3; Sequence=VSP_003225;
CC Name=4; Synonyms=delE214E4;
CC IsoId=Q30201-4; Sequence=VSP_003219, VSP_003225;
CC Name=5;
CC IsoId=Q30201-5; Sequence=VSP_003219;
CC Name=6;
CC IsoId=Q30201-6; Sequence=VSP_003220;
CC Name=7; Synonyms=delE3;
CC IsoId=Q30201-7; Sequence=VSP_003221;
CC Name=8; Synonyms=1043-2283del, intron6ins;
CC IsoId=Q30201-8; Sequence=VSP_003226, VSP_003227;
CC Name=9; Synonyms=delE3-7;
CC IsoId=Q30201-9; Sequence=VSP_003223, VSP_003224;
CC Name=10; Synonyms=562-878del;
CC IsoId=Q30201-10; Sequence=VSP_003222;
CC -!- TISSUE SPECIFICITY: IN ALL TISSUES TESTED EXCEPT BRAIN.
CC -!- DISEASE: DEFECTS IN HFE ARE A CAUSE OF HEREDITARY HEMOCHROMATOSIS
CC (HH). HH IS AN AUTOSOMAL RECESSIVE INBORN DISORDER OF IRON
CC METABOLISM. FREQUENT AMONG CAUCASIANS. HH IS CHARACTERIZED BY
CC ABNORMAL INTESTINAL IRON ABSORPTION AND PROGRESSIVE INCREASE OF
CC TOTAL BODY IRON, WHICH RESULTS IN MIDLIFE IN CLINICAL
CC COMPLICATIONS INCLUDING CIRRHOSIS, CARDIOPATHY, DIABETES,
CC ENDOCRINE DYSFUNCTIONS, ARTHROPATHY, AND SUSCEPTIBILITY TO LIVER
CC CANCER. SINCE THE DISEASE COMPLICATIONS CAN BE EFFECTIVELY
CC PREVENTED BY REGULAR PHLEBOTOMIES, EARLY DIAGNOSIS IS MOST
CC IMPORTANT TO PROVIDE A NORMAL LIFE EXPECTANCY TO THE AFFECTED
CC SUBJECTS.
CC -!- DISEASE: DEFECTS IN HFE ARE A CAUSE OF PORPHYRIA CUTANEA TARDA
CC (PCT), A DISORDER CHARACTERIZED BY LIGHT-SENSITIVE DERMATITIS AND
CC PRESENCE OF LARGE AMOUNTS OF UROPORPHYRIN IN URINE. IRON OVERLOAD
CC IS OFTEN PRESENT IN ASSOCIATION WITH VARYING DEGREES OF LIVER
CC DAMAGE. PCT IS THE MOST COMMON FORM OF PORPHYRIA WORLDWIDE. IT
CC OCCURS IN TWO FORMS: THE SPORADIC TYPE (PCT TYPE I) AND THE
CC FAMILIAL TYPE (PCT TYPE II), WHICH IS INHERITED IN AN AUTOSOMAL

Query Match 99.2%; Score 1502; DB 1; Length 348;
 Best Local Similarity 99.3%; Pred. No. 2,36-118;
 Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDLGLSFEALGYVDDQLFVYDHSRRVPRTPWSSRISQ 60
 DB 23 RLLRSHSLHYLFMGASQDLGLSFEALGYVDDQLFVYDHSRRVPRTPWSSRISQ 82

QY 61 MWLQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
 DB 83 MWLQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 142

QY 121 QDALEFCPDLTDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
 DB 143 QDHLFCPDLTDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202

QY 181 DQVPPPLVKKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKVLPNGDG 240
 DB 203 DQVPPPLVKKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKVLPNGDG 262

QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
 DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 2

HFE DICSU STANDARD; PRT; 348 AA.

AC QSLG42;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hereditary hemochromatosis protein precursor.
 GN HFE.

OS Dicerorhinus sumatrensis (Sumatran rhinoceros).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Dicerorhinus.
 OX NCBI_TaxID=89632;
 [1]
 RN SEQUENCE FROM N.A.
 RA West C.J., Worley M., Beutler E.;
 RT "Rhinoceros HFE polymorphisms."
 RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 CC affinity for iron-loaded transferrin.
 CC -!- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -!- SIMILARITY: TO MHC CLASS I ANTIGENS.

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 CC
 CC EMBL; AY007543; AAG23703.1; -
 CC HSSP; Q30201; 1A6Z.
 CC InterPro; IPR007110; Ig-like.
 CC InterPro; IPR003597; Ig_c1.
 CC InterPro; IPR003006; Ig_MHC.
 CC InterPro; IPR001039; MHC_I.
 CC Pfam; PF00047; Ig_1.
 CC Pfam; PF00129; MHC_I; 1.
 CC ProDom; PD000050; MHC_I; 1.
 CC SMART; SM00407; Igc1; 1.
 CC PROSITE; PSS0835; IG_LIKE; 1.
 CC PROSITE; PS00290; IG_MHC; 1.
 CC MHC_I; Transmembrane; Glycoprotein; Signal.
 KW MHC_I; 22 BY SIMILARITY.
 FT SIGNAL 1 22
 FT CHAIN 23 348 HEREDITARY HEMOCHROMATOSIS PROTEIN.
 FT DOMAIN 23 114 EXTRACELLULAR ALPHA-1.

FT DOMAIN 115 205 EXTRACELLULAR ALPHA-2.
 FT DOMAIN 206 297 EXTRACELLULAR ALPHA-3.
 FT TRANSMEM 307 330 CONNECTING PEPTIDE.
 FT TRANSMEM 331 348 POTENTIAL.
 FT DISULFID 124 187 CYTOPLASMIC TAIL.
 FT DISULFID 225 282 BY SIMILARITY.
 FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 130 130 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 234 234 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 348 AA; 39740 MW; 518BFD357AB83B90 CRC64;

Query Match 81.0%; Score 1227; DB 1; Length 348;
 Best Local Similarity 81.0%; Pred. No. 2,2e-95;
 Matches 221; Conservative 20; Mismatches 32; Indels 0; Gaps 0;

QY 4 RSHSLHYLFMGASQDLGLSFEALGYVDDQLFVYDHSRRVPRTPWSSRISQMWL 63
 DB 26 RSHSLHYLFMGASQDLGLSFEALGYVDDQLFVYDHSRRVPRTPWSSRISQMWL 85

QY 64 QLSQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDGQA 123
 DB 86 QLSQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDGQH 145

QY 124 LDFCPDPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVLDOQ 183
 DB 146 LDFCPDPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVLDOQ 205

QY 184 VPPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKVLPNGDGTQY 243
 DB 206 VPPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKVLPNGDGTQY 265

QY 244 GWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
 DB 266 SWALAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 3

HFE CERSI STANDARD; PRT; 348 AA.

AC Q9GKZ0;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hereditary hemochromatosis protein precursor.
 GN HFE.

OS Ceratotherium simum (White rhinoceros) (Square-lipped rhinoceros).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Ceratotherium.
 OX NCBI_TaxID=9807;
 [1]
 RN SEQUENCE FROM N.A.
 RA West C.J., Worley M., Beutler E.;
 RT "Rhinoceros HFE polymorphisms."
 RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 CC affinity for iron-loaded transferrin.
 CC -!- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -!- SIMILARITY: TO MHC CLASS I ANTIGENS.

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 CC or send an email to license@isb-sib.ch).
 CC
 CC EMBL; AY007541; AAG23701.1; -
 CC HSSP; Q30201; 1A6Z.
 CC InterPro; IPR007110; Ig-like.
 CC InterPro; IPR003597; Ig_c1.
 CC InterPro; IPR003006; Ig_MHC.
 CC InterPro; IPR001039; MHC_I.
 CC Pfam; PF00047; Ig_1.
 CC Pfam; PF00129; MHC_I; 1.
 CC ProDom; PD000050; MHC_I; 1.
 CC SMART; SM00407; Igc1; 1.
 CC PROSITE; PSS0835; IG_LIKE; 1.
 CC PROSITE; PS00290; IG_MHC; 1.
 CC MHC_I; Transmembrane; Glycoprotein; Signal.
 KW MHC_I; 22 BY SIMILARITY.
 FT SIGNAL 1 22
 FT CHAIN 23 348 HEREDITARY HEMOCHROMATOSIS PROTEIN.
 FT DOMAIN 23 114 EXTRACELLULAR ALPHA-1.

DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR PRINTS; PR01638; MHCCLASSI.
 DR PRODOM; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGC1; 1.
 DR DR PROSITE; PS00835; IG LIKE; 1.
 DR DR PROSITE; PS00290; IG_MHC; 1.
 KW MHC I; Transmembrane; Glycoprotein; Signal.
 FT SIGNAL 1 22 BY SIMILARITY.
 FT CHAIN 23 348 HEREDITARY HEMOCHROMATOSIS PROTEIN.
 FT DOMAIN 23 115 EXTRACELLULAR ALPHA-1.
 FT DOMAIN 115 205 EXTRACELLULAR ALPHA-2.
 FT DOMAIN 206 297 EXTRACELLULAR ALPHA-3.
 FT DOMAIN 298 306 CONNECTING PEPTIDE.
 FT TRANSMEM 307 330 POTENTIAL.
 FT DOMAIN 331 348 CYTOPLASMIC TAIL.
 FT DISULFID 124 187 BY SIMILARITY.
 FT DISULFID 225 282 BY SIMILARITY.
 FT CARBOHYD 110 130 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 130 130 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 234 234 N-LINKED (GLCNAC. .) (POTENTIAL).
 SQ SEQUENCE 348 AA; 39822 MW; 2523016EC9FBE91 CRC64;

Query Match 80.9%; Score 1225; DB 1; Length 348;
 Best Local Similarity 81.3%; Pred. No. 3.2e-95;
 Matches 222; Conservative 18; Mismatches 33; Indels 0; Gaps 0;

QY 4 RSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRVETPTWVSSRISSQMWL 63
 |||||
 DB 26 RSHSLRYLFMGASERDGLPLFEALGYVDDQLFVFDHESRRVETPTWVSSRISSQMWL 85
 |||||

QY 64 QLSQSLKGDWDMFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGDGDA 123
 |||||
 DB 86 QLSQSLKGDWDMFTVDFWTIMDNHNHSHKESHTLQVILGCEVQEDNSTGRGFWKYGVDGDH 145
 |||||

QY 124 LEFCPTDLDWRAAEPRAPWTKLEWRHKIRARQNRAYLERDCPAQLQQLLELGRGVLDQ 183
 |||||
 DB 146 LEFCPETLDWRAAESRALTTKLEWVKNIRAKQNRAYLERDCPEQLQWLELGRGVLDQ 205
 |||||

QY 184 VPPLVKVTHVTSVTLRCALNYFPQNTMKWLKDKQPMDAKEPEPKDVLNPGDGTQ 243
 |||||
 DB 206 VPPLVKVTHVASAVTLRCQALNFPQNTMRWLKDKRPMDVDAESKDVLPSGDGTQ 265
 |||||

QY 244 GWITLAVPPGEQRYTCQVEHPGLDQPLIVIE 276
 |||||
 DB 266 SWEALAVPPGEQRYTCQVEHPGLDQPLTATWE 298
 |||||

RESULT 4
 HFE_RHTUN
 ID_HFE_RHTUN STANDARD; PRT; 348 AA.
 AC Q9GL41; 2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hereditary hemochromatosis protein precursor.
 GN HFE.
 OS Rhinoceros unicornis (Greater Indian rhinoceros).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Rhinoceros.
 OX NCBI_TaxID=9809;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA West C.J., Worley M., Beutler E.;
 RT "Rhinoceros HFE polymorphisms";
 RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
 CC -|- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 CC affinity for iron-loaded transferrin.
 CC -|- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -|- SIMILARITY: TO MHC CLASS I ANTIGENS.

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DR EMBL; AY007544; AAG23704.1; -
 DR HSP; Q30201; IAGZ.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; IG_C1.
 DR InterPro; IPR003006; IG_MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR PRINTS; PR01638; MHCCLASSI.
 DR PRODOM; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGC1; 1.
 DR DR PROSITE; PS00835; IG LIKE; 1.
 DR DR PROSITE; PS00290; IG_MHC; 1.
 KW MHC I; Transmembrane; Glycoprotein; Signal.
 FT SIGNAL 1 22 BY SIMILARITY.
 FT CHAIN 23 348 HEREDITARY HEMOCHROMATOSIS PROTEIN.
 FT DOMAIN 23 114 EXTRACELLULAR ALPHA-1.
 FT DOMAIN 115 205 EXTRACELLULAR ALPHA-2.
 FT DOMAIN 206 297 EXTRACELLULAR ALPHA-3.
 FT DOMAIN 298 306 CONNECTING PEPTIDE.
 FT TRANSMEM 307 330 POTENTIAL.
 FT DOMAIN 331 348 CYTOPLASMIC TAIL.
 FT DISULFID 124 187 BY SIMILARITY.
 FT DISULFID 225 282 BY SIMILARITY.
 FT CARBOHYD 110 130 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 130 130 N-LINKED (GLCNAC. .) (POTENTIAL).
 FT CARBOHYD 234 234 N-LINKED (GLCNAC. .) (POTENTIAL).
 SQ SEQUENCE 348 AA; 39743 MW; F2723D57A327A6B4 CRC64;

Query Match 80.6%; Score 1221; DB 1; Length 348;
 Best Local Similarity 80.6%; Pred. No. 7e-95;
 Matches 220; Conservative 20; Mismatches 33; Indels 0; Gaps 0;

QY 4 RSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRVETPTWVSSRISSQMWL 63
 |||||
 DB 26 RSHSLRYLFMGASERDGLPLFEALGYVDDQLFVFDHESRRVETPTWVSSRISSQMWL 85
 |||||

QY 64 QLSQSLKGDWDMFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGDGDA 123
 |||||
 DB 86 QLSQSLKGDWDMFTVDFWTIMDNHNHSHKESHTLQVILGCEVQEDNSTGRGFWKYGVDGDH 145
 |||||

QY 124 LEFCPTDLDWRAAEPRAPWTKLEWRHKIRARQNRAYLERDCPAQLQQLLELGRGVLDQ 183
 |||||
 DB 146 LEFCPETLDWRAAESRALTTKLEWVKNIRAKQNRAYLERDCPEQLQWLELGRGVLDQ 205
 |||||

QY 184 VPPLVKVTHVTSVTLRCALNYFPQNTMKWLKDKQPMDAKEPEPKDVLNPGDGTQ 243
 |||||
 DB 206 VPPLVKVTHVASAVTLRCQALNFPQNTMRWLKDKRPMDVDAESKDVLPSGDGTQ 265
 |||||

QY 244 GWITLAVPPGEQRYTCQVEHPGLDQPLIVIE 276
 |||||
 DB 266 SWEALAVPPGEQRYTCQVEHPGLDQPLTATWE 298
 |||||

RESULT 5

HFE_DICBI
 ID_HFE_DICBI STANDARD; PRT; 348 AA.
 AC Q9GL43; Q9GK81;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hereditary hemochromatosis protein precursor.
 GN HFE.
 OS Dicerus bicornis (Black rhinoceros).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Rhinoceros.

OX NCBI_TaxID=9805;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA West C.J., Worley M., Beutler E.;
 RT "Rhinceros HFE polymorphisms";
 RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 CC affinity for iron-loaded transferrin.
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- SIMILARITY: TO MHC CLASS I ANTIGENS.
 CC -----
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 CC -----
 DR EMBL; AY007542; AAG23702.1; -;
 DR EMBL; AF301592; AAG39940.1; -;
 DR HSSP; Q30201; 1A6Z.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig_c1.
 DR InterPro; IPR003006; Ig_MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; Ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR ProDom; PD000050; MHC_I; 1.
 DR SMART; SM00407; Igc1; 1.
 DR PROSITE; PS50835; IG_LIKE; 1.
 DR PROSITE; PS00230; IG_MHC; 1.
 DR MHC I; Transmembrane; Glycoprotein; Signal.
 FT SIGNAL 1 22 BY SIMILARITY.
 FT CHAIN 23 348 HEREDITARY HEMOCHROMATOSIS PROTEIN.
 FT DOMAIN 115 205 EXTRACELLULAR ALPHA-1.
 FT DOMAIN 206 297 EXTRACELLULAR ALPHA-2.
 FT DOMAIN 298 306 EXTRACELLULAR ALPHA-3.
 FT TRANSMEM 307 330 CONNECTING PEPTIDE.
 FT DOMAIN 331 348 POTENTIAL.
 FT DISULFID 124 187 CYTOPLASMIC TAIL.
 FT BY SIMILARITY.
 FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 130 130 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 234 234 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 348 AA; 39818 MW; 4D95E7B01E48FB90 CRC64;
 Query Match 80.4%; Score 1218; DB 1; Length 348;
 Best Local Similarity 80.6%; Pred. No. 1.2e-94;
 Matches 220; Conservative 20; Mismatches 33; Indels 0; Gaps 0;
 QY 4 RSHSLHYLFNGASEQDLGLSLFALGVDDQLFVFDHESRRVPEPTPWSSRISQMWL 63
 DB 26 RSHSLRYLFNGASERDGLFLFALGVDDQLFVFDHESRRVPEPTPWSSRISQMWL 85
 QY 64 QLSQSLKGDWDMFTVDFWTIMENHNASKESHTLQVILGCEMQRDSTEGYWKYGYDGQDA 123
 DB 86 QLTQSLKGDWDMFTVDFWTIMDNHNHNSKESHTLQVILGCEVQBDNSTRGFWKYGYDGDH 145
 QY 124 LEFCPTDLWRAAEPRAPWPTKLEWERHKIRARQNAYLERDCPAQLQLLELGRVLDQ 183
 DB 146 LEFCPTDLWRAAESRALTTKLEWEVKNIRAKQNAYLERDCPEQLQLLELGRVLDQ 205
 QY 184 VPPLVKVTHVTSVTLRCLALNYFPQNTMKWLKDKQPMDAKEPDKVDLPNGDGTQ 243
 DB 206 VPPLVKVTHVTSVTLRCLALNYFPQNTMKWLKDKPVDVKAESKDVLPNGDGTQ 265
 QY 244 GWITLAVPPGEORYTCVHEHGLDQPLIIVE 276
 DB 266 SWEALAVPPGEORYTCVHEHGLDQPLTATWE 298

RESULT 6
 HFE RAT
 ID HFE RAT STANDARD; PRT; 360 AA.
 AC O35799; O35175;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hereditary hemochromatosis protein homolog precursor (RTI-CAFE).
 GN HFE.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OC NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Banasch M.W., Schaefer H., Schmidt W.E.;
 RL Submitted (SEP-1997) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE OF 42-303 FROM N.A.
 RC TISSUE=Small intestine;
 RA Sawada-Hirai R., Rothenberg B.E.;
 RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: BINDS TO TRANSFERRIN RECEPTOR (TFR) AND REDUCES ITS
 CC AFFINITY FOR IRON-LOADED TRANSFERRIN (BY SIMILARITY).
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- SIMILARITY: TO MHC CLASS I ANTIGENS.
 CC -----
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 CC -----
 DR EMBL; AJ001517; CAA04799.1; -;
 DR EMBL; AF008587; AAB86597.1; -;
 DR HSSP; Q30201; 1A6Z.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig_c1.
 DR InterPro; IPR003006; Ig_MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; Ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR PRINTS; PR01638; MHCCLASSI.
 DR ProDom; PD000050; MHC_I; 1.
 DR SMART; SM00407; Igc1; 1.
 DR PROSITE; PS50835; IG_LIKE; 1.
 DR PROSITE; PS00230; IG_MHC; 1.
 DR MHC I; Transmembrane; Glycoprotein; Signal.
 FT SIGNAL 1 25 POTENTIAL.
 FT CHAIN 26 360 HEREDITARY HEMOCHROMATOSIS PROTEIN
 FT HOMOLOG.
 FT DOMAIN 26 127 EXTRACELLULAR ALPHA-1.
 FT DOMAIN 128 218 EXTRACELLULAR ALPHA-2.
 FT DOMAIN 219 310 EXTRACELLULAR ALPHA-3.
 FT DOMAIN 311 319 CONNECTING PEPTIDE.
 FT TRANSMEM 320 340 POTENTIAL.
 FT DOMAIN 341 360 CYTOPLASMIC TAIL.
 FT BY SIMILARITY.
 FT DISULFID 137 200 BY SIMILARITY.
 FT CARBOHYD 115 115 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 143 143 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 167 167 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 247 247 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CONFLICT 198 198 R -> K (IN REF. 2).
 SQ SEQUENCE 360 AA; 40988 MW; CC819834EB3240B3 CRC64;
 Query Match 75.6%; Score 1145; DB 1; Length 360;
 Best Local Similarity 73.2%; Pred. No. 1.6e-88;
 Matches 205; Conservative 29; Mismatches 38; Indels 8; Gaps 1;

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Qy 5 SHSLHYLFMGASEODLGLSFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQWLQ 64
Db 32 SHSLHYLFMGASEODLGLSFEALGYVDDQLFVSYNHESRAEPRAPIWILGOTSSQLWLQ 91
Qy 65 LSQSLKGDWDMFTVDFTWIMENHNASK-----ESHYLOVILGCEMQEDNSTEGYWKY 116
Db 92 LSQSLKGDWDMFTVDFTWIMENHNASK-----ESHYLOVILGCEMQEDNSTEGYWKY 151
Qy 117 GYGQDQALEFCPDTLDWRAAEPRAPWPKLEWERHKIRARONRAYLERDCPAQLQQLLELG 176
Db 152 GYGQDQHLFCPKTLNWSAAEPRAWATMEWEHRIRAROSRYLDQDCPQQLQVLELQ 211
Qy 177 RGVLDDQVPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEFEKDVLP 236
Db 212 RGVLDDQVPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEFEKDVLP 271
Qy 237 NGDGTGQWTLAVPQGEORYTCQVHPGLDQPLVIWE 276
Db 272 NGDGTGQWTLAVPQGEORYTCQVHPGLDQPLVIWE 311

RESULT 7
HFE_MOUSE STANDARD; PRT; 359 AA.
AC P70387;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hereditary hemochromatosis protein homolog precursor.
GN HFE OR MR2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RN SEQUENCE FROM N.A.
RC STRAIN=129/SvJ;
RC MDLINE=98060831; PubMed=9396865;
RA Riegert P., Gilfillan S., Nanda I., Schmid M., Bahram S.;
RT "The mouse HFE gene.";
RL Immunogenetics 47:174-177 (1998).
RN [2]
RN SEQUENCE FROM N.A.
RC STRAIN=BALB/c; TISSUE=Lung;
RA Hashimoto K.;
RL Submitted (SEP-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RN SEQUENCE OF 37-211 FROM N.A.
RC STRAIN=BALB/c; TISSUE=Liver;
RC MDLINE=97148566; PubMed=9020055;
RA Hashimoto K., Hirai M., Kurosawa Y.;
RT "Identification of a mouse homolog for the human hereditary
hemochromatosis candidate gene.";
RL Biochem. Biophys. Res. Commun. 230:35-39 (1997).
RN [4]
RN SEQUENCE OF 79-359 FROM N.A.
RC STRAIN=129;
RA Albright W., Drabant B., Doenecke D.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
CC -I- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
affinity for iron-loaded transferrin (By similarity).
CC -I- SUBCELLULAR LOCATION: Type I membrane protein.
CC -I- SIMILARITY: TO MHC CLASS I ANTIGENS.
CC -----
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CC -----
EMBL; AF007558; AAC03447.1; -.
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DR EMBL; U66849; AAB07525.1; -
DR EMBL; Y12650; CAA73197.1; -
DR EMBL; U80604; AAB51504.1; -
DR PIR; JC5382; JC5382.
DR HSSP; Q30201; IA62.
DR MGD; MGI:109191; Hfe.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_c1.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGc1; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 359
FT HEREDITARY HEMOCHROMATOSIS PROTEIN
FT HOMOLOG.
FT DOMAIN 25 126
FT DOMAIN 127 217
FT DOMAIN 218 309
FT DOMAIN 310 318
FT TRANSMEM 319 339
FT DOMAIN 340 359
FT DISULFID 136 139
FT DISULFID 237 294
FT CARBOHYD 114 114
FT CARBOHYD 142 142
FT CARBOHYD 166 166
FT CARBOHYD 246 246
FT SEQUENCE 359 AA; 40548 MW; 4BDE6C27F9FF20B4 CRC64;
SQ
Query Match 74.6%; Score 1129; DB 1; Length 359;
Best Local Similarity 71.9%; Pred. No. 3.5e-87;
Matches 202; Conservative 30; Mismatches 41; Indels 8; Gaps 1;
Qy 4 RSHSLHYLFMGASEODLGLSFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQWL 63
Db 30 RSHSLHYLFMGASEODLGLSFEALGYVDDQLFVSYNHESRAEPRAPIWILGOTSSQLWL 89
Qy 64 QLSQSLKGDWDMFTVDFTWIMENHNASK-----ESHYLOVILGCEMQEDNSTEGYWK 115
Db 90 HLSQSLKGDWDMFTVDFTWIMENHNASK-----ESHYLOVILGCEMQEDNSTEGYWK 149
Qy 116 YGYDQDQALEFCPDTLDWRAAEPRAPWPKLEWERHKIRARONRAYLERDCPAQLQQLLEL 175
Db 150 YGYDQDQALEFCPDTLDWRAAEPRAPWPKLEWERHKIRARONRAYLERDCPAQLQQLLEL 209
Qy 176 GRGVLDQVPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEFEKDVLP 235
Db 210 GRGVLDQVPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEFEKDVLP 269
Qy 236 PNGDGTGQWTLAVPQGEORYTCQVHPGLDQPLVIWE 276
Db 270 PNGDGTGQWTLAVPQGEORYTCQVHPGLDQPLVIWE 310

RESULT 8
HFE_MOUSE STANDARD; PRT; 361 AA.
AC P01894;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE RLA class I histocompatibility antigen, alpha chain 11/11 precursor.
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxID=9986;
RN [1]
```

RP SEQUENCE FROM N.A.
RX MEDLINE=84290724; PubMed=6432910;
RA Tykocinski M.L., Marche P.N., Max E.E., Kindt T.J.;
RT "Rabbit class I MHC genes: cDNA clones define full-length transcripts
of an expressed gene and a putative pseudogene.";
RL J. Immunol. 133:2261-2269(1984).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
MICROGLOBULIN).
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or send an email to license@isb-sib.ch).
CC
CC EMBL; K02441; AAA98729.1; -.
DR PIR; A02193; HLBB.
DR HSP; Q30201; IA6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig ci.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
DR MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 361 RLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT ALPHA CHAIN 1/11.
FT DOMAIN 25 114 EXTRACELLULAR ALPHA-1.
FT DOMAIN 115 206 EXTRACELLULAR ALPHA-2.
FT DOMAIN 207 298 EXTRACELLULAR ALPHA-3.
FT DOMAIN 299 308 CONNECTING PEPTIDE.
FT TRANSMEM 309 329 CYTOPLASMIC.
FT DOMAIN 330 361 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT DISULFID 125 188 BY SIMILARITY.
FT DISULFID 227 283 BY SIMILARITY.
SQ SEQUENCE 361 AA; 580B673323C1AE35 CRC64;
Query Match 34.1%; Score 517; DB 1; Length 361;
Best Local Similarity 40.1%; Pred. No. 4.9e-36;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;
QY 5 SHSLHYLFMGASQDLGLSLFEALGYVDDQLFPVYDHE--SRVPRTPWVSSRISQW 62
DB 26 SHSMRYFTYSVRPGLEPRFIIVGVDDTQFVRFDSDAASPRMEQAPWM-QGVPEY 84
QY 63 LQLSQLKGDHMFVTVDFTWMENHNASK-SHTLQVILGCMEQDNS-TEGYWYGYDG 120
DB 85 DQQTQIAKDTAQTFRNLNTALRYNQSAAGSHFTQTFMFCVWADGRPHGYQYADG 144
QY 121 QDALEFCPTDLWRAPAEPRAMPYKLEWERHKBARONRAYLERDCPAQLQOLLEGRV 180
DB 145 ADYIALNEDLRSWTAADTAQNTQKWEAAG-EAERHRAVLERECVLEWLRYLEMGKET 203
QY 181 DQOVPLVYKTHVTSS-VTTLCRALNYYPQNTMKWLKDKQPMADKPEPDKVLNPGD 239
DB 204 QRADPPRAHVTHHPASDREATHLCWALGFYPASISLTWQDGDSD-QTQDTVELTRPGD 262
QY 240 GTYQGMITLAVPPGEQRYTCOVHEPGLDQPLIVWE 276
DB 263 GTQKWAADVVPSEGEQRYTCRVQHEGLPEPLITWE 299

RESULT 9
HAIB_RABIT STANDARD; PRT; 361 AA.
ID HAIB_RABIT
AC P06140;
DT 01-JAN-1988 (Rel. 06, Created)
DT 01-JAN-1988 (Rel. 06, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE RLA class I histocompatibility antigen, alpha chain 19-1 precursor.
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_Taxid:9986;
RN [1]
SEQUENCE FROM N.A.
RP MEDLINE=85103547; PubMed=3917974;
RA Marche P.N., Tykocinski M.L., Max E.E., Kindt T.J.;
RT "Structure of a functional rabbit class I MHC gene: similarity to
human class I genes.";
RL Immunogenetics 21:71-82(1985).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
MICROGLOBULIN).
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CC
CC EMBL; K02819; AAA98730.1; -.
DR PIR; I46858; I46858.
DR HSP; Q30201; IA6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig ci.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
DR MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 361 RLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT ALPHA CHAIN 19-1.
FT DOMAIN 25 114 EXTRACELLULAR ALPHA-1.
FT DOMAIN 115 206 EXTRACELLULAR ALPHA-2.
FT DOMAIN 207 298 EXTRACELLULAR ALPHA-3.
FT DOMAIN 299 308 CONNECTING PEPTIDE.
FT TRANSMEM 309 329 CYTOPLASMIC.
FT DOMAIN 330 361 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT DISULFID 125 188 BY SIMILARITY.
FT DISULFID 227 283 BY SIMILARITY.
SQ SEQUENCE 361 AA; 40455 MW; C06FBD8B87ED0546 CRC64;
Query Match 34.1%; Score 517; DB 1; Length 361;
Best Local Similarity 40.1%; Pred. No. 4.9e-36;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;
QY 5 SHSLHYLFMGASQDLGLSLFEALGYVDDQLFPVYDHE--SRVPRTPWVSSRISQW 62
DB 26 SHSMRYFTYSVRPGLEPRFIIVGVDDTQFVRFDSDAASPRMEQAPWM-QGVPEY 84
QY 63 LQLSQLKGDHMFVTVDFTWMENHNASK-SHTLQVILGCMEQDNS-TEGYWYGYDG 120
DB 85 DQQTQIAKDTAQTFRNLNTALRYNQSAAGSHFTQTFMFCVWADGRPHGYQYADG 144

QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWHRKIRARONRAYLERDCPAQLQQLLELGRGVL 180
 Db 145 ADYIALNEDLRSTWTAADTAQNTQKWEAAG-EAERHAYLERECVWELRYLENGKETL 203
 QY 181 DQVPPPLVKVTHVTS-VTTLRCRALNYYPQNTMKWLKDKQPMDAKEFEKPKDVLNPGD 239
 Db 204 QRADPPKRAVTHHPASDREATLRCAWALGFYPAEISLTWQDGED-QTQDTVELVETRPAGD 262
 QY 240 GTYQGWITLAVPGEORVTCOVHEPGLDPLIVINE 276
 Db 263 GTQKMAAVVVPGEORVTCOVHEPGLDPLIVINE 299

RESULT 10

LA01 PANTR ID LA01 PANTR STANDARD; PRT; 365 AA.
 AC PI6209;
 DT 01-APR-1990 (Rel. 14, Created)
 DT 01-APR-1990 (Rel. 14, Last sequence update)
 DT 01-APR-1993 (Rel. 25, Last annotation update)
 DE CHLA class I histocompatibility antigen, A-2 alpha chain precursor.
 DE Pan troglodytes (Chimpanzee).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
 OX NCBI_TaxID=9598;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=90201944; PubMed=1690682;
 RA Lawlor D.A., Warren E., Ward F.E., Parham P.;
 RT "Comparison of class I MHC alleles in humans and apes.";
 RL Immunol. Rev. 113:147-185(1990).
 CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
 CC THE IMMUNE SYSTEM.
 CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
 CC MICROGLOBULIN).
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 CC or send an email to license@isb-sib.ch).
 CC EMBL; M30678; AAA7970.1; --
 DR PIR; I36961; I36961.
 DR HSP; Q95352; IHKK.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig cl.
 DR InterPro; IPR003006; Ig_MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; Ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR PRINTS; PR01638; MHCCLASS1.
 DR ProDom; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGC1; 1.
 DR PROSITE; PS50835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 DR MHC I; Transmembrane; Glycoprotein; Signal.
 KW SIGNAL 1 24
 FT CHAIN 25 365 CHLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
 FT FT 25 365 A-2 ALPHA CHAIN.
 FT DOMAIN 25 114 EXTRACELLULAR ALPHA-1.
 FT DOMAIN 115 206 EXTRACELLULAR ALPHA-2.
 FT DOMAIN 207 298 EXTRACELLULAR ALPHA-3.
 FT DOMAIN 299 308 CONNECTING PEPTIDE.
 FT TRANSMEM 309 332
 FT DOMAIN 333 365 CYTOPLASMIC TAIL.
 FT DISULFID 125 188 BY SIMILARITY.
 FT DISULFID 227 283 BY SIMILARITY.
 FT CARBOHYD 110 110 N-LINKED (GLCNAC...) (BY SIMILARITY).
 SQ SEQUENCE 365 AA; 40848 MW; FC452786BD038D3E CRC64;

Query Match 33.8%; Score 511; DB 1; Length 365;
 Best Local Similarity 39.7%; Pred. No. 1.6e-35;
 Matches 110; Conservative 44; Mismatches 115; Indels 8; Gaps 7;
 QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHE--SRVPEPTPWSSRISQMW 62
 Db 26 SHSMRYFTTSVSPGCGEPRTIAVGVDVDTQVRFDSDAASQRMPEPRAPWIEQE-GPEYW 84
 QY 63 LQLSLSLKGDWHMFTVDFTWIMENNASKE-SHTLQVILGCEMQEDNS-TEGYWYGYDGD 120
 Db 85 DEETSAKASQTDVRDVLGTLRGYINQSDGSHITQIMYGCDVSDGRFLRGTRQDAYDG 144
 QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWHRKIRARONRAYLERDCPAQLQQLLELGRGVL 180
 Db 145 ADYIALNEDLRSTWTAADTAQNTQKWEAAG-EAERHAYLERECVWELRYLENGKETL 203
 QY 181 DQVPPPLVKVTHH-VTSSVTTLRCRALNYYPQNTMKWLKDKQPMDAKEFEKPKDVLNPGD 239
 Db 204 QRTDPPKTHHTHPISDHEATLRCAWALGFYPAEISLTWQDGED-QTQDTVELVETRPAGD 262
 QY 240 GTYQGWITLAVPGEORVTCOVHEPGLDPLIVINE 276
 Db 263 GTQKMAAVVVPGEORVTCOVHEPGLDPLIVINE 299
 RESULT 11
 HAIB BOVIN ID HAIB BOVIN STANDARD; PRT; 364 AA.
 AC PI3753;
 DT 01-JAN-1990 (Rel. 13, Created)
 DT 01-JAN-1990 (Rel. 13, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE BOLA class I histocompatibility antigen, alpha chain BL3-7 precursor.
 OS Bos taurus (Bovine).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
 OX NCBI_TaxID=9913;
 RN [1]
 RP SEQUENCE FROM N.A.
 RX MEDLINE=88258075; PubMed=3133413;
 RA Ennis P.D., Jackson A.P., Parham P.;
 RT "Molecular cloning of bovine class I MHC cDNA.";
 RL J. Immunol. 141:642-651(1988).
 CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
 CC THE IMMUNE SYSTEM.
 CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
 CC MICROGLOBULIN).
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 CC or send an email to license@isb-sib.ch).
 CC EMBL; M21043; AAA30641.1; --
 DR HSP; PI6391; IED3.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig cl.
 DR InterPro; IPR003006; Ig_MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; Ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR PRINTS; PR01638; MHCCLASS1.
 DR ProDom; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGC1; 1.
 DR PROSITE; PS50835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 KW MHC I; Transmembrane; Glycoprotein; Signal.
 FT SIGNAL 1 27
 FT CHAIN 28 364 BOLA CLASS I HISTOCOMPATIBILITY ANTIGEN,

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FT DOMAIN 28 117 ALPHA CHAIN BL3-7.
FT DOMAIN 118 209 EXTRACELLULAR ALPHA-1.
FT DOMAIN 210 301 EXTRACELLULAR ALPHA-2.
FT DOMAIN 302 310 EXTRACELLULAR ALPHA-3.
FT TRANSMEM 311 331 CONNECTING PEPTIDE.
FT DOMAIN 332 364 CYTOPLASMIC.
FT CARBOHYD 106 106 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 113 113 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT DISULFID 128 191 BY SIMILARITY.
FT DISULFID 230 286 BY SIMILARITY.
SQ SEQUENCE 364 AA; 41513 MW; 622056CF7DCFF7873 CRC64;

Query Match 33.7%; Score 510; DB 1; Length 364;
Best Local Similarity 38.9%; Pred. No. 1.9e-35;
Matches 109; Conservative 49; Mismatches 114; Indels 8; Gaps 7;

QY 2 LLRSHSLHVLPMGASEODLGLSLFEALGYVDQLFVYDHE--SRRVERPTWVSSRISS 59
DB 26 LAGSHSLRYPTGYVSRPLGEPRIAVGYVDDTQFVRFSDAPNPREPRVPMWQEGP 84
QY 60 QMWLQLSLSLKGWDHMTVDFTWIMENHNASKE-SHTLQVILGCEMOEDNS-TEGYWKYG 117
DB 85 EYDWRNTRIYKDTAIFRVDLNTLRGYNQSGTSHNIQAMYCDVGPDCRLLRGFWQFG 144
QY 118 YDQDLEFCDFDLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGR 177
DB 145 YDGRDVIATNEELRSWTAADTAQAITKRWAAAG-AAETWRNLYEGECVWELRRYLENGK 203
QY 178 GVLDQVQVPLVKTTH-VTSVTLTLCRALNYQNTWKLXDKQPMDAKEPEPKDVL 236
DB 204 DTLRLADPPKAVHTHSISDREVTLCRWALGFYPEISLWQREGD-QTDWELVETRP 262
QY 237 NGDGTQGWITLAVPGEQRYTCQVEHPLGLDPLIWI 276
DB 263 SGGDTQKWAALVPSGEQRYTCVQHGSLQPLRLWE 302

RESULT 12
ID 1A11 HUMAN STANDARD; PRT; 365 AA.
AC P13746;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE HLA class I histocompatibility antigen, A-11 alpha chain precursor.
GN HLA-A OR HLA-A.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (A*1101/A*1102).
RX MEDLINE=89030641; PubMed=2460344;
RA Mayer W.E., Jonker M., Klein D., Ivanyi P., van Seventer G.,
RA Klein J.;
RT "Nucleotide sequences of chimpanzee MHC class I alleles: evidence for
RT trans-species mode of evolution.";
RL EMBO J. 7:2765-2774(1988).
RN [2]
RP SEQUENCE FROM N.A. (A*1101/A*1102).
RX MEDLINE=94287401; PubMed=8016845;
RA Lin L., Tokunaga K., Ishikawa Y., Bannai M., Kashiwase K.,
RA Kuwata S., Akaza T., Tadokoro K., Shibata Y., Juji T.;
RT "Sequence analysis of serological HLA-A11 split antigens, A11.1 and
RT A11.2.";
RL Tissue Antigens 43:78-82(1994).
RN [3]
RP SEQUENCE OF 26-365 FROM N.A. (A*1101).
RX MEDLINE=87192928; PubMed=2437024;
RA Cowan E.P., Jelachich M.L., Biddison W.E., Coligan J.E.;
RT "DNA sequence of HLA-A11: remarkable homology with HLA-A3 allows
RT identification of residues involved in epitopes recognized by

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RT antibodies and T cells.";
RL Immunogenetics 25:241-250(1987).
CC -I- FUNCTION: Involved in the presentation of foreign antigens to
CC the immune system.
CC -I- SUBUNIT: Dimer of alpha chain and a beta chain (beta-2-
CC microglobulin).
CC -I- POLYMORPHISM: THE FOLLOWING ALLELES OF A-11 ARE KNOWN: A*1101 (A-
CC 11E) AND A*1102 (A-11K). THE SEQUENCE SHOWN IS THAT OF A*1101.
CC -----
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; X13111; CAA31503.1; --
CC EMBL; X13112; CAA31504.1; --
CC EMBL; D16841; BAA04117.1; --
CC EMBL; D16842; BAA04118.1; --
CC EMBL; M16010; AAA65449.1; --
CC EMBL; M16007; AAA65449.1; JOINED.
CC EMBL; M16008; AAA65449.1; JOINED.
CC EMBL; M16009; AAA65449.1; JOINED.
CC PIR; I83063; I83063.
CC PIR; S03536; A47636.
CC HSSP; O19673; 1HSB.
CC MIM; 142800; --
CC GO; GO:0005887; C:integral to plasma membrane; NAS.
CC GO; GO:0030106; F:MHC class I receptor activity; NAS.
CC GO; GO:0006955; P:immune response; NAS.
CC InterPro; IPR007110; IG-like.
CC InterPro; IPR003597; IG_c1.
CC InterPro; IPR003006; IG_MHC.
CC InterPro; IPR001039; MHC_I.
CC Pfam; PF00047; IG; 1.
CC Pfam; PF00129; MHC_I; 1.
CC PRINTS; PR01638; MHCCLASSI.
CC ProDom; PD000050; MHC_I; 1.
CC SMART; SM00407; IGc1_1.
CC PROSITE; PS00835; IG-LIKE; 1.
CC PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal; Polymorphism.
FT SIGNAL 1 24
FT CHAIN 25 365 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT A-11 ALPHA CHAIN.
FT DOMAIN 25 114 EXTRACELLULAR ALPHA-1.
FT DOMAIN 115 206 EXTRACELLULAR ALPHA-2.
FT DOMAIN 207 298 EXTRACELLULAR ALPHA-3.
FT DOMAIN 299 308 CONNECTING PEPTIDE.
FT TRANSMEM 309 332
FT DOMAIN 333 365 CYTOPLASMIC TAIL.
FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT DISULFID 125 188 BY SIMILARITY.
FT DISULFID 227 283 BY SIMILARITY.
FT VARIANT 43 43 E -> K (IN ALLELE A*1102).
FT /FTID=VAR 004353.
SQ SEQUENCE 365 AA; 40937 MW; FE449CE2D4BF6CC5 CRC64;

Query Match 33.6%; Score 508; DB 1; Length 365;
Best Local Similarity 39.4%; Pred. No. 2.8e-35;
Matches 109; Conservative 46; Mismatches 114; Indels 8; Gaps 7;

QY 5 SHSLHVLPMGASEODLGLSLFEALGYVDQLFVYDHE--SRRVERPTWVSSRISSQW 62
DB 26 SHSMRYFTSVSRPGRGEPRFAVGYVDVTQVRFSDAASQRMPEAPWIEQE-GPEYW 84
QY 63 LOLSLSLKGWDHMTVDFTWIMENHNASKE-SHTLQVILGCEMOEDNS-TEGYWKYG 120
DB 85 DQETRNKQAQSDTRDVLGTLRGYNOQSGDGHITQIMYGCDVGPDCRLLRGYRDADYG 144
QY 121 QDALEFCDFDLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGR 180

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Db 145 KDVIALLNEDLSRTAADMAAQITKRWAAH-AAEQRAYLEGRCVWELRYLNGKETL 203
QY 181 DQOVPLPVKVTTH-VTSVVTTLRCALNYPQNTMKWLKDKQPMADAKEFEPKDVLPNGD 239
Db 204 QRTDPKTHMTHTPIDSHETALRCWALGFYPAEITLTWQDGED-QTQDTVELVETRPAGD 262
QY 240 GTYQGMITLAVPGEQRYTCQVEHPLGLDPLIVWE 276
Db 263 GTFQKMAAVVPSGEGORYTCHVQHEGLPKPLTLRWE 299

RESULT 13
ID 1A03 HUMAN STANDARD; PRT; 370 AA.
AC P04439;
DT 13-AUG-1987 (Rel. 05, Created)
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE HLA class I histocompatibility antigen, A-3 alpha chain precursor.
GN HLA-A OR HLA*
OS Homo sapiens (Human)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN 1
RP SEQUENCE FROM N.A. (A*0301).
RX MEDLINE=84207948; PubMed=6609814;
RA Strachan T., Sodoyer R., Damotte M., Jordan B.R.;
RT "Complete nucleotide sequence of a functional class I HLA gene,
RT HLA-A3: implications for the evolution of HLA genes.";
RL EMBO J. 3:887-894 (1984).
RN 2
RP SEQUENCE FROM N.A. (A*0302).
RX MEDLINE=85290871; PubMed=2993417;
RA Cowan E.P., Jordan B.E., Coligan J.E.;
RT "Molecular cloning and DNA sequence analysis of genes encoding
RT cytotoxic T lymphocyte-defined HLA-A3 subtypes: the E1 subtype.";
RL J. Immunol. 135:2835-2841 (1985).
CC -1- FUNCTION: Involved in the presentation of foreign antigens to
CC the immune system.
CC -1- SUBUNIT: Dimer of alpha chain and a beta chain (beta-2-
CC microglobulin).
CC -1- POLYMORPHISM: THE FOLLOWING ALLELES OF A-3 ARE KNOWN: A*0301 (A-
CC 3.1) AND A*0302. THE SEQUENCE SHOWN IS THAT OF A*0301.
CC -----
CC This SWISS-PROT entry is copyright. It is produced through a collaboration
CC between the Swiss Institute of Bioinformatics and the EMBL outstation -
CC the European Bioinformatics Institute. There are no restrictions on its
CC use by non-profit institutions as long as its content is in no way
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CC or send an email to license@isb-sib.ch).
CC -----
CC EMBL; X00492; CAA2162.1; ALT_TERM.
DR PIR; A02192; HLHUA3.
DR HSSP; O19673; IHSB.
DR MTM; 142800;
DR GO; GO:0005887; C:integral to plasma membrane; NAS.
DR GO; GO:0030106; F:MHC class I receptor activity; NAS.
DR GO; GO:0006955; P:immune response; NAS.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_1.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal; Polymorphism.

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FT SIGNAL 1 29
FT CHAIN 30 370
FT FT HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT A-3 ALPHA CHAIN
FT FT EXTRACELLULAR ALPHA-1.
FT DOMAIN 30 119
FT DOMAIN 120 211
FT DOMAIN 212 303
FT DOMAIN 304 313
FT TRANSMEM 314 337
FT DOMAIN 338 370
FT CARBOHYD 115 115
FT DISULFID 130 193
FT DISULFID 232 288
FT VARIANT 181 181
FT VARIANT 185 185
FT SEQUENCE 370 AA; 41368 MW; ABBIFA77460318A2 CRC64;
SQ
Query Match 33.4%; Score 506; DB 1; Length 370;
Best Local Similarity 39.6%; Pred. No. 4.2e-35;
Matches 110; Conservative 46; Mismatches 112; Indels 10; Gaps 8;

QY 5 SHSLHYLFMGASEQDGLSLFALGVDDQLPVFDHE--SRVEPRTFWSSRSQMW 62
Db 31 SHSMRYFTTSVSRPGRGEPRTIAGVYDDTQFVRFDSDAASQRMPEAPWIEQ-GPEYW 89
QY 63 LQLSQSLKGDHMTFTVDFTIMENNASKE-SHTLQVLGCEMQEDNS-TEGYWKYGYDG 120
Db 90 DQETRNVAQSQSTDRVDLTGLTGYNQSEAGSHITQIMYGCVDGSDGRFLGRTRQDAYDG 149
QY 121 QDALSPCPDPTLDWRAAEPRAPWTKLEWE--RHKIRARQNRAYLERDCPAQLQLELGRGV 179
Db 150 KDVIALLNEDLSRTAADMAAQITKRWAAH-AAEQRAYLDGTCEVLELRYLNGKET 207
QY 180 LQOVVPLPVKVTTH-VTSVVTTLRCALNYPQNTMKWLKDKQPMADAKEFEPKDVLPNG 238
Db 208 LQRTDPPKTHMTHTPIDSHETALRCWALGFYPAEITLTWQDGED-QTQDTVELVETRPAG 266
QY 239 DGTQGMITLAVPGEQRYTCQVEHPLGLDPLIVWE 276
Db 267 DGTQKMAAVVPSGEGORYTCHVQHEGLPKPLTLRWE 304

RESULT 14
ID 1A80 HUMAN STANDARD; PRT; 365 AA.
AC Q09160;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE HLA class I histocompatibility antigen, AW-80(A-1) alpha chain
DE precursor.
GN HLA-A OR HLA*
OS Homo sapiens (Human)
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN 1
RP SEQUENCE FROM N.A.
RX MEDLINE=94245293; PubMed=8188325;
RA Balas A., Garcia-Sanchez F., Gomez-Reino F., Vicario J.L.;
RT "Characterization of a new and highly distinguishable HLA-A allele in
RT a Spanish family";
RL Immunogenetics 39:452-452 (1994).
RN 2
RP SEQUENCE FROM N.A.
RA Domena J.D.;
RL Submitted (JUN-1993) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
CC -1- POLYMORPHISM: THE ONLY ALLELE OF AW-80 KNOWN IS A*8001 WHICH IS

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DR MIM; 142800; --
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig-cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal; Polymorphism.
FT SIGNAL 1 24
FT CHAIN 25 365
FT
FT HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT A-31 ALPHA CHAIN.
FT DOMAIN 25 114
FT DOMAIN 115 206
FT DOMAIN 207 298
FT DOMAIN 299 308
FT DOMAIN 309 332
FT TRANSMEM 333 365
FT DOMAIN 110 110
FT CARBOHYD 125 188
FT DISULFID 227 283
FT VARIANT 121 121
FT
FT M -> I (IN ALLELE A*3104).
FT /FTid=VAR_010373.
FT Q -> R (IN ALLELE A*3104).
FT /FTid=VAR_010374.
FT
FT VARIANT 138 138
FT
SQ SEQUENCE 365 AA; 41004 MW; 4E760C821A3C553B CRC64;

Query Match 33.2%; Score 502; DB 1; Length 365;
Best Local Similarity 39.0%; Pred. No. 8.9e-35;
Matches 108; Conservative 50; Mismatches 111; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDLGLSLFEALGYVDQLFVFDHE--SRVEPRTPWVSSRISSQMW 62
   |||:| | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 26 SHSMRYFTTSVRGGEPRFIAVGIVDDTQFVRFDSDAASQRMFPAPWIEQE-RPEYW 84
   |:::| | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 63 LQLSOSLKGWDHMTVDFTWIMENHNASKS-SHTLQVILGCENQEDNS-TEGYWKYGYDG 120
   |:::| | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 85 DQETRNKAKHSQIDRVDLGLTGLRGYNQSEAGSHTIQMNYGCDVGSDFLRGYQDDAYDG 144
   |:::| | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 121 QDALEFCPDLDNRAEPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
   :| | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 145 KDYIALNEDLRSTADMAAQITQRKEAARV-AEQLRAYLEGTCTVWELRYLENGKETL 203
   :| | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 181 DQVPPPLVKVTHH-VTSSVTTLCRALNYYPNITMKWLKDKQPMDAKEFEPKDVLPNGD 239
   :| | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 204 QRTDPPKTHMTHAVSDHEATLSCWALSFPYPAEITLTWQRDGED-QTQDTLVELVETRPAGD 262
   :| | | | | | | | | | | | | | | | | | | | | | | | | | | |
QY 240 GTYQGWITLAVPGEORPTCVQHPGLDQPLVIWE 276
   ||:| | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 263 GTFQKASVVVPSGQEQRYTCHVQHEGLPKPLTRWE 299
   ||:| | | | | | | | | | | | | | | | | | | | | | | | | | | |
```

Search completed: August 5, 2003, 13:08:49
Job time : 10.5 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:05:29 ; Search time 32 Seconds
(without alignments)

2225.704 Million cell updates/sec

Title: US-10-092-404-3

Perfect score: 1514

Sequence: 1 RLRSLSLHLYFMGASEQDL.....RYTCQVEHGLDQPLIVWE 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL_23:*
1: sp_archaea:*
2: sp_bacteria:*
3: sp_fungi:*
4: sp_human:*
5: sp_invertebrate:*
6: sp_mammal:*
7: sp_mhc:*
8: sp_organelle:*
9: sp_phase:*
10: sp_plant:*
11: sp_rodent:*
12: sp_virus:*
13: sp_vertebrate:*
14: sp_unclassified:*
15: sp_rvirus:*
16: sp_bacteriap:*
17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1129	74.6	358	11 Q8C2A6	Q8c2a6 mus musculus
2	1129	74.6	359	11 Q9D754	Q9d754 mus musculus
3	792	52.3	272	11 Q9R105	Q9r105 rattus norv
4	574	37.9	116	4 Q9HC69	Q9hc69 homo sapien
5	540.5	35.7	359	7 Q8HX81	Q8hx81 ornithorhyn
6	537.5	35.5	354	7 Q95HB3	Q95hb3 anas platyr
7	537.5	35.1	340	7 Q9BD50	Q9bd50 pongo pygma
8	530.5	35.0	334	7 Q9BQK3	Q9bqk3 homo sapien
9	530.5	35.0	341	4 Q9NPL2	Q9npl2 homo sapien
10	530.5	35.0	341	7 Q9B460	Q9b460 homo sapien
11	530.5	35.0	341	7 Q9BCU3	Q9bcu3 pan troglod
12	527.5	34.8	341	7 Q9BCU4	Q9bcu4 pan troglod
13	515	34.0	356	7 Q8HX66	Q8hx66 sus scrofa
14	514	33.9	332	7 Q3O990	Q3o990 pan troglod
15	514	33.9	365	7 Q9TPL7	Q9tpl7 pan troglod
16	512	33.8	105	4 Q9HC71	Q9hc71 homo sapien

ALIGNMENTS

RESULT 1

Q8C2A6 PRELIMINARY; PRT; 358 AA.

ID Q8C2A6;
AC Q8C2A6;
DT 01-WAR-2003 (Tremblrel. 23, Created)
DT 01-WAR-2003 (Tremblrel. 23, Last sequence update)
DT 01-WAR-2003 (Tremblrel. 23, Last annotation update)
DE Hemochromatosis.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]_TaxID=10090;
RP SEQUENCE FROM N.A.
RC STRAIN=NOD; TISSUE=Thymus;
RX MEDLINE=22354683; PubMed=12466851;
RA The FANTOM Consortium,
RA the RIKEN Genome Exploration Research Group Phase I & II Team;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs."
RL Nature 420:563-573 (2002).
DR EMBL; AK088986; BAC40688.1; -.
SQ SEQUENCE 358 AA; 40421 MW; EE88FB6E5AAC844D CRC64;

Query Match 74.6%; Score 1129; DB 11; Length 358;
Best Local Similarity 71.9%; Pred. No. 1.3e-97;
Matches 202; Conservative 30; Mismatches 41; Indels 8; Gaps 1;

QY 4 RSHSLHLYFMGASEQDLGLSLFEALGYVDDQLFVFDHESRVERVEPTPMVSRISQMWL 63
Db 29 RSHSLHLYFMGASEQDLGLSLFEALGYVDDQLFVFDHESRVERVEPTPMVSRISQMWL 88
QY 64 QLSQSLKGDHMTFTVDFWTIMENHNASK-----ESHTLVQILGCENQDNSTGYWK 115
Db 89 HLSQSLKGDWYMFVDFWTIMGNYNHVKYKLGVVVSESHILQVLGCEVHEDNSTSGFWR 148
QY 116 YGYDQDALLEFCDFDLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQLEL 175
Db 149 YGYDQDHLFCFCKTLNWSAEPGAWATKVEWDEHKIRAKQNRDYLEKDCPQLKRLLEL 208

QY	176	GRGVLDQQVPLPKVTHVTSTVTLRCRALNYPQNI	TMKWLKDQKPMDAKEFEFKDVL	235
		GRGVLDQQVPLPKVTHVTSTVTLRCRALNYPQNI	TMKWLKDQKPMDAKEFEFKDVL	
DB	209	GRGVLGQVPTLVKVTTRHWAGTSLRQALDFFPQNI	TMRLKDNQDPLDAKDVNPEKVL	268
		GRGVLGQVPTLVKVTTRHWAGTSLRQALDFFPQNI	TMRLKDNQDPLDAKDVNPEKVL	
QY	236	PNGDGTQGWITFLAVPPGEEQRYTCQVEHPGLDPLIVWE	276	
		PNGDGTQGWITFLAVPPGEEQRYTCQVEHPGLDPLIVWE	276	
DB	269	PNGDGTQGWITFLAVAPDGTFRFCQVEHPGLDPLTASWE	309	
		PNGDGTQGWITFLAVAPDGTFRFCQVEHPGLDPLTASWE	309	

QY 126 FCDTLDRAAEPRAPWTKLEWRHKIRARQNRAYLERDCPAQLQQLLELGRGVLDQV 185
 DB 144 LDKDTWTTFAADAAQITKKKEEDGTVAERKYYLLENTCIEWLRKYRYGKDVLERRR 203
 QY 186 PLVKVTHHTVSSVTTLCRALNYYPQNTMKWLKQKPMDAKEFEPRKDVLPNGDGTG 245
 DB 204 PEVRSGMEADKILSLSCRAHGYPPIPSISWLKGM-VQEQTQSGSTVPNSDGIYHIW 262
 QY 246 ITLAVPGBEQRVTCQVHPLGLOPLIVWE 276
 DB 263 ATIDVVPGRKDKYQCRVHASLPQGLFSWE 293

RESULT 7
 Q9BD50 PRELIMINARY; PRT; 340 AA.
 ID Q9BD50
 AC Q9BD50
 DT 01-JUN-2001 (T-EMBLrel. 17, Created)
 DT 01-JUN-2001 (T-EMBLrel. 17, Last sequence update)
 DT 01-MAR-2003 (T-EMBLrel. 23, Last annotation update)
 DE MHC class I related protein MRL isoform.
 GN MRL.
 OS Pongo pygmaeus (Orangutan).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pongo.
 OX NCBI_TaxID=9600;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Placenta;
 RX MEDLINE=99003494; PubMed=9784382;
 RA Yamaguchi H., Kurosawa Y., Hashimoto K.;
 RT "Expanded genomic organization of conserved mammalian MHC class I-
 related genes, human MRL and its murine ortholog.";
 RL Biochem. Biophys. Res. Commun. 250:558-564(1998).
 CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
 CC IMMUNE SYSTEM (BY SIMILARITY).
 CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
 CC MICROGLOBULIN) (BY SIMILARITY).
 DR EMBL; AJ271828; CAC28215.1; -.
 DR HSSP; Q30201; 1A6Z.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig cl.
 DR InterPro; IPR003006; Ig MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; Ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR ProDom; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGcl; 1.
 DR PROSITE; PS00835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 KW Glycoprotein; Transmembrane.
 FT NON_TER 1
 SQ SEQUENCE 340 AA; 39375 MW; A893952B78725F17 CRC64;

Query Match 35.1%; Score 531.5; DB 7; Length 340;
 Best Local Similarity 39.1%; Pred. No. 1.4e-41;
 Matches 106; Conservative 51; Mismatches 111; Indels 3; Gaps 3;
 QY 4 RSHSLHYLFWGASEQDGLSLFALGYVDQDLFVFDHESRRVPRTPVSSRISQMWL 63
 DB 23 RTHSLRYFLRGVSDPIHGVPFISGVYDSHPITTYDSVTROKEPRAPMAENLADPHWE 82
 QY 64 QLSQSLKGDHMTVDFTWTTMENHNASKESHTLQVLGCMQBDNSTEGYWKYGYDQDA 123
 DB 83 RYTQLLRGQOMKVELKRLQRHYNHSGSHYQRMIGCELLEDGSGTTGFLQYAYDQDF 141
 QY 124 LEFCPDTLDWRAAEPRAPWTKLEWRHKIRARQNRAYLERDCPAQLQQLLELGRGVLDQ 183
 DB 142 LIENKDTLSLWADVNDVAHTIKRAWEANQHELOQKNWLEBECIAWLKRFLEYGKDTLQRT 201
 QY 184 VPLVKVTHHTVSSVTTLCRALNYYPQNTMKWLKQKPMDAKEFEPRKDVLPNGDGT 242
 DB 202 EPLVRVNRKRTFPFGVTALFCAHGFYPPEIYNTWMNGEEI-VQEMDYGDILPDSGDGT 260
 QY 243 QGMITLAVPGBEQRVTCQVHPLGLOPLIV 273

DB 261 QTWASFELDPQSSNLYSCHVEHGVHMLQV 291
 RESULT 8
 Q9TK3 PRELIMINARY; PRT; 334 AA.
 ID Q9TK3
 AC Q9TK3
 DT 01-MAY-2000 (T-EMBLrel. 13, Created)
 DT 01-MAY-2000 (T-EMBLrel. 13, Last sequence update)
 DT 01-MAR-2003 (T-EMBLrel. 23, Last annotation update)
 DE MHC class I-related protein MRL (Fragment).
 GN MRL.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OX NCBI_TaxID=9606;
 RN [1]
 RP SEQUENCE FROM N.A.
 RC TISSUE=Placenta;
 RX MEDLINE=99003494; PubMed=9784382;
 RA Yamaguchi H., Kurosawa Y., Hashimoto K.;
 RT "Expanded genomic organization of conserved mammalian MHC class I-
 related genes, human MRL and its murine ortholog.";
 RL Biochem. Biophys. Res. Commun. 250:558-564(1998).
 CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
 CC IMMUNE SYSTEM (BY SIMILARITY).
 CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
 CC MICROGLOBULIN) (BY SIMILARITY).
 DR EMBL; AF073485; AAC72900.1; -.
 DR EMBL; AF073484; AAC72900.1; JOINED.
 DR HSSP; Q30201; 1A6Z.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig cl.
 DR InterPro; IPR003006; Ig MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; Ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR ProDom; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGcl; 1.
 DR PROSITE; PS00835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 KW Glycoprotein; Transmembrane.
 FT NON_TER 1
 SQ SEQUENCE 334 AA; 38586 MW; 4C3E3A8248A39BA4 CRC64;
 Query Match 35.0%; Score 530.5; DB 7; Length 334;
 Best Local Similarity 39.1%; Pred. No. 1.6e-41;
 Matches 106; Conservative 50; Mismatches 112; Indels 3; Gaps 3;
 QY 4 RSHSLHYLFWGASEQDGLSLFALGYVDQDLFVFDHESRRVPRTPVSSRISQMWL 63
 DB 16 RTHSLRYFLRGVSDPIHGVPFISGVYDSHPITTYDSVTROKEPRAPMAENLADPHWE 75
 QY 64 QLSQSLKGDHMTVDFTWTTMENHNASKESHTLQVLGCMQBDNSTEGYWKYGYDQDA 123
 DB 76 RYTQLLRGQOMKVELKRLQRHYNHSGSHYQRMIGCELLEDGSGTTGFLQYAYDQDF 134
 QY 124 LEFCPDTLDWRAAEPRAPWTKLEWRHKIRARQNRAYLERDCPAQLQQLLELGRGVLDQ 183
 DB 135 LIENKDTLSLWADVNDVAHTIKRAWEANQHELOQKNWLEBECIAWLKRFLEYGKDTLQRT 194
 QY 184 VPLVKVTHHTVSSVTTLCRALNYYPQNTMKWLKQKPMDAKEFEPRKDVLPNGDGT 242
 DB 195 EPLVRVNRKRTFPFGVTALFCAHGFYPPEIYNTWMNGEEI-VQSIDYGDILPDSGDGT 253
 QY 243 QGMITLAVPGBEQRVTCQVHPLGLOPLIV 273
 DB 254 QAWASIELDPQSSNLYSCHVEHGVHMLQV 284
 RESULT 9
 Q9NPL2

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ID Q9NPL2 PRELIMINARY; PRT; 341 AA.
AC Q9NPL2;
DT 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Mrl protein.
GN Mrl.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=peripheral blood;
RX MEDLINE=20470599; PubMed=11019920;
RA Parra-Cuadrado J.F., Navarro P., Mirones I., Setien F., Oteo M.,
RA Martinez-Naves E.;
RT "A study on the polymorphism of human MHC class I-related Mrl gene and
RT identification of an Mrl-like pseudogene.";
RL Tissue Antigens 56:170-172(2000).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; AJ249778; CAB77667.1; -.
DR HSSP; Q30201; 1A6Z.
DR InterPro; IPR007110; Ig-like.
DR PRINTS; PR01638; MHCCLASSI.
DR PRODOM; PD000050; MHC I; 1.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRODOM; PD000050; MHC_I; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
DR Glycoprotein; Transmembrane.
KW Glycoprotein; Transmembrane.
SQ SEQUENCE 341 AA; 39366 MW; 2990C1F3F0A1CAD9 CRC64;

Query Match 35.0%; Score 530.5; DB 4; Length 341;
Best Local Similarity 39.1%; Pred. No. 1.7e-41;
Matches 106; Conservative 50; Mismatches 112; Indels 3; Gaps 3;

QY 4 RSHSLHYLFMGASEQDLGLSLFALGYVDQDLFFVYDHSRRVPRTPWSSRISSQMWL 63
Db 23 RTHSLRYFLRGVSDPIHGVPFISVGVDSPHTTYSVTQKEPRAPWAENLADPHWE 82
QY 64 QLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMQBDNSTEGYWKYGYDGQDA 123
Db 83 RYTQLLRGWQOMFKVELKRLQRYNHS -GSHTYQRMIGCELLEDGSTTGLQYAYDQDF 141
QY 124 LEFCPTDLWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 142 LIENKDTLSLWADVNAHTIKQAEANQHLLYQKNWLEECIAWLKRFLEYKCDTLQRT 201
QY 184 VPLVAVKTHVT -SSVTLTLCRALNYYPQNTMKWLKDKQPMDAKEFEKPVLPNGDGT 242
Db 202 EPLVVRNKRKTFPGVTALFCKAHGFYPPPIYTWKNGEEI -VQIDYGDILLPSDGT 260
QY 243 QGMITLAVPPGEQRVTCQVHEPGLDQPLIV 273
Db 261 QAWASIELDPQSSNLYSCHVEHCGVHMVLQV 291

RESULT 10
Q95460 PRELIMINARY; PRT; 341 AA.
AC Q95460;
DT 01-FEB-1997 (TrEMBLrel. 02, Created)
DT 01-FEB-1997 (TrEMBLrel. 02, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Class I histocompatibility antigen-like protein.

ID Q9BCU3 PRELIMINARY; PRT; 341 AA.
AC Q9BCU3;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE MHC class I related protein, Mr1B1 isoform.
GN Mr1.
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RA Martinez-Naves E.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.

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OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=thymus;
RX MEDLINE=95350662; PubMed=7624800;
RA Hashimoto K., Hirai M., Kurosawa Y.;
RT "A gene outside the human MHC related to classical HLA class I
RT genes.";
RL Science 269:693-695(1995).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; U22963; AAC50174.1; -.
DR HSSP; Q30201; 1A6Z.
DR Genew; HGNC:4975; HLALS.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003006; Ig MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRODOM; PD000050; MHC_I; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
SQ SEQUENCE 341 AA; 39366 MW; 2990C1F3F0A1CAD9 CRC64;

Query Match 35.0%; Score 530.5; DB 7; Length 341;
Best Local Similarity 39.1%; Pred. No. 1.7e-41;
Matches 106; Conservative 50; Mismatches 112; Indels 3; Gaps 3;

QY 4 RSHSLHYLFMGASEQDLGLSLFALGYVDQDLFFVYDHSRRVPRTPWSSRISSQMWL 63
Db 23 RTHSLRYFLRGVSDPIHGVPFISVGVDSPHTTYSVTQKEPRAPWAENLADPHWE 82
QY 64 QLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMQBDNSTEGYWKYGYDGQDA 123
Db 83 RYTQLLRGWQOMFKVELKRLQRYNHS -GSHTYQRMIGCELLEDGSTTGLQYAYDQDF 141
QY 124 LEFCPTDLWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 142 LIENKDTLSLWADVNAHTIKQAEANQHLLYQKNWLEECIAWLKRFLEYKCDTLQRT 201
QY 184 VPLVAVKTHVT -SSVTLTLCRALNYYPQNTMKWLKDKQPMDAKEFEKPVLPNGDGT 242
Db 202 EPLVVRNKRKTFPGVTALFCKAHGFYPPPIYTWKNGEEI -VQIDYGDILLPSDGT 260
QY 243 QGMITLAVPPGEQRVTCQVHEPGLDQPLIV 273
Db 261 QAWASIELDPQSSNLYSCHVEHCGVHMVLQV 291

RESULT 11
Q9BCU3 PRELIMINARY; PRT; 341 AA.
AC Q9BCU3;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE MHC class I related protein, Mr1B1 isoform.
GN Mr1.
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RA Martinez-Naves E.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.

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RP SEQUENCE FROM N.A.
RA Parra-Cuadrado J.F., Garcia-Pavia P., Gomez del Moral M.;
RT "Identification of MRL cDNA sequences in non-human primates";
RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
IMMUNE SYSTEM (BY SIMILARITY).
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
MICROGLOBULIN) (BY SIMILARITY).
CC EMBL; AJ275984; CAC34272.1; --
DR HSSP; Q30201; 1A6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; IG MHC; 1.
DR Glycoprotein; Transmembrane.
KW Glycoprotein; Transmembrane.
FT VARIANT 197 197
SQ SEQUENCE 341 AA; 33394 MW; FBF822BCAB2C7A8 CRC64;
Query Match 35.0%; Score 530.5; DB 7; Length 341;
Best Local Similarity 39.1%; Pred. No. 1.7e-41;
Matches 106; Conservative 51; Mismatches 111; Indels 3; Gaps 3;
QY 4 RSHSLHYLFMGASEQDLGLSLFALGVYDDQLFVFDHESRRVETPTWVSSRISSQMWL 63
DB 23 RTHSLRYFRGLGSDPIHGVPFISVGYVDSHPITTYDSVTROKEPRAPMAENLAPDHW 82
QY 64 QLSQSLKGDHMTVDFTWIMENHNASKESHHTLQVILGCEMQEDNSTEGYWKYGYDQDA 123
DB 83 RYTQLRGWQMKVELKRLQRYNHS -GSHTYQRMIGCELLEDGSTTGFQVAYDQGF 141
QY 124 LEFCPTLDWRAAEPRAPWTKLEWRHKIRARQNRAYLERDCPAQLQQLLELGRGVLDQ 183
DB 142 LIENKDTLSLWADVNDVAHTIKQAEANQHLLYQKNWLEEECIAWLKRFLYEGKDTLQRT 201
QY 184 VPLVKVTHVT-SSVTLTLCRALNYYPQNIITKWLKDKQPMDAKEPEFKDVLPGDGT 242
DB 202 EPLVVRNKRKTFPGVTALFCKAHGFYPPEIYMTWMKNGEEI-VQSIDYGDILPSGDGT 260
QY 243 QGWITLAVPGEQRYTCQVHPGLDQPLIV 273
DB 261 QTWASVELDPQSSNLISYCHVCHGVHVLQV 291
RESULT 12
Q9BCU4 PRELIMINARY; PRT; 341 AA.
AC Q9BCU4
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE MHC class I related protein, MR1B1 isoform.
GN MR1.
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RA Martinez-Naves E.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
RN [2]
CC SEQUENCE FROM N.A.
RA Parra-Cuadrado J.F., Garcia-Pavia P., Gomez del Moral M.;
RT "Identification of MRL cDNA sequences in non-human primates";
RL Submitted (MAR-2001) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
IMMUNE SYSTEM (BY SIMILARITY).

CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
MICROGLOBULIN) (BY SIMILARITY).
CC EMBL; AJ275982; CAC34274.1; --
DR HSSP; Q30201; 1A6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; IG MHC; 1.
KW Glycoprotein; Transmembrane.
FT VARIANT 197 197
SQ SEQUENCE 341 AA; 33382 MW; DFF16AF1FAB2D272 CRC64;
Query Match 34.8%; Score 527.5; DB 7; Length 341;
Best Local Similarity 39.1%; Pred. No. 3.2e-41;
Matches 106; Conservative 50; Mismatches 112; Indels 3; Gaps 3;
QY 4 RSHSLHYLFMGASEQDLGLSLFALGVYDDQLFVFDHESRRVETPTWVSSRISSQMWL 63
DB 23 RTHSLRYFRGLGSDPIHGVPFISVGYVDSHPITTYDSVTROKEPRAPMAENLAPDHW 82
QY 64 QLSQSLKGDHMTVDFTWIMENHNASKESHHTLQVILGCEMQEDNSTEGYWKYGYDQDA 123
DB 83 RYTQLRGWQMKVELKRLQRYNHS -GSHTYQRMIGCELLEDGSTTGFQVAYDQGF 141
QY 124 LEFCPTLDWRAAEPRAPWTKLEWRHKIRARQNRAYLERDCPAQLQQLLELGRGVLDQ 183
DB 142 LIENKDTLSLWADVNDVAHTIKQAEANQHLLYQKNWLEEECIAWLKRFLYEGKDTLQRT 201
QY 184 VPLVKVTHVT-SSVTLTLCRALNYYPQNIITKWLKDKQPMDAKEPEFKDVLPGDGT 242
DB 202 EPLVVRNKRKTFPGVTALFCKAHGFYPPEIYMTWMKNGEEI-VQSIDYGDILPSGDGT 260
QY 243 QGWITLAVPGEQRYTCQVHPGLDQPLIV 273
DB 261 QTWASVELDPQSSNLISYCHVCHGVHVLQV 291
RESULT 13
Q8HX66 PRELIMINARY; PRT; 356 AA.
AC Q8HX66
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE MHC class I antigen (Fragment).
GN SLA-1.
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suidae; Sus.
OX NCBI_TaxID=9823;
RN [1]
RP SEQUENCE FROM N.A.
RA Martens G.W., Baker J.E., Smith D.M.;
RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY135589; AAN35107.1; --
FT NON_TER 1
SQ SEQUENCE 356 AA; 39585 MW; 94FC7A461DBF555B CRC64;
Query Match 34.0%; Score 515; DB 7; Length 356;
Best Local Similarity 39.9%; Pred. No. 5.1e-40;
Matches 110; Conservative 47; Mismatches 111; Indels 8; Gaps 7;
QY 6 HSLHYLFMGASEQDLGLSLFALGVYDDQLFVFDHESRRVETPTWVSSRISSQMWL 63
DB 19 HSLRYFYTAVSRPDLGDSRFYAVGVDDTQFVRPDSADPNPMEPRAPWQIQE-QGEYWD 77
QY 64 QLSQSLKGDHMTVDFTWIMENHNASKESHHTLQVILGCEMQEDNSTEGYWKYGYDQ 121

Db 78 ETRNMGSAQNDKRVLDLTKLGRYNGSEAGSHITQRMVGYDGDGLLLGRYDQDAYDGA 137
QY 122 DALEFCFDDTLDRAAEPRAPWPTKLEWERHKIRARQNAYLERDCPAQLQQLLELGRGVLD 181
Db 138 DYIALNEDLRSWTAADTAQAQITKKEWAAV-AEQERSYLEGTCVWQLQKYLEMGKDTLQ 196
QY 182 QOVPPPLVKVTHVTSV-TTLRCALNYPONITMKWLKQKQMDAKFEPKQVLPNGDG 240
Db 197 RAEPFKTHVRHPSFSDGLVTLGRWALGFYKPEISLTWQREGQD-QSQDMELVETPSGDG 255
QY 241 TYQGVITLAVPPGGEQRYTCOVHPGLDQPLIVWE 276
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AC Q30990;
DT 01-NOV-1996 (Tremblrel. 01, Created)
DT 01-NOV-1996 (Tremblrel. 01, Last sequence update)
DT 01-MAR-2003 (Tremblrel. 23, Last annotation update)
DE Chimpanzee MHC class I Ch1a chain (Fragment).
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=89235215; PubMed=2715640;
RA Parham P., Lawlor D.A., Lomen C.E., Ennis P.D.;
RT "Diversity and diversification of HLA-A,B,C alleles.";
RL J. Immunol. 142:3937-3950(1989).
CC -I- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC -I- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
CC EMBL; M24047; AAA35426.1; -.
DR HSSP; Q95352; 1HHK.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_c1.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
FT NON TER 332
SQ SEQUENCE 332 AA; 37433 MW; 9AA9A55DF979360 CRC64;

Query Match 33.9%; Score 514; DB 7; Length 332;
Best Local Similarity 40.1%; Pred. No. 5.8e-40;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;
QY 5 SHSLHFLPMGASQDGLSLFEALGYVDOLFFVFDHE--SRVPRTPWVSRRSSQW 62
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QY 63 LQLSQSLKGDHMTVDFTWIMENHNASKS-SHTLQVILGCEMQEDNS-TEGYWKYGYDG 120
Db 85 DQETRAKSAHSQTDRLDGLTGRYNGSEAGSHITQIMYGCVDGSDGRFLRGYRDQAYDG 144
QY 121 QDALEFCFDDTLDRAAEPRAPWPTKLEWERHKIRARQNAYLERDCPAQLQQLLELGRGVLD 180
Db 145 KDVIALLNEDLRSWTAADTAQAQITKKEWAAH-AAEQRAYLEGTCTVWELRRYLENGKETL 203
QY 181 DQOVPPPLVKVTHH-VTSVSTTLRCALNYPONITMKWLKQKQMDAKFEPKQVLPNGDG 239

Db 204 QRTDPPKTHMTHHPISDHEATRLCNALGFYPAEITLTWQDGED-QTQDTLTVETRPAGD 262
QY 240 GTYQGVITLAVPPGGEQRYTCOVHPGLDQPLIVWE 276
Db 263 GTFOKWAALVPPGGEQRYTCOVHPGLDQPLIVWE 299
RESULT 15
Q9TPL7
ID Q9TPL7 PRELIMINARY; PRT; 365 AA.
AC Q9TPL7;
DT 01-MAY-2000 (Tremblrel. 13, Created)
DT 01-MAY-2000 (Tremblrel. 13, Last sequence update)
DT 01-MAR-2003 (Tremblrel. 23, Last annotation update)
DE MHC class I antigen.
GN HLA-A OR PATR-A.
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=35A-1;
RX MEDLINE=99335357; PubMed=10405321;
RA Matsui M., Machida S., Feinstein S.M., Akatsuka T.;
RT "Molecular analyses of five new chimpanzee MHC class I alleles:
RT Implications for differences between evolutionary mechanisms of HLA-A,
RT -B, and -C loci.";
RL Biochem. Biophys. Res. Commun. 261:46-52(1999).
RN [2]
RP SEQUENCE FROM N.A.
RX TISSUE=Blood;
RX MEDLINE=20322475; PubMed=10866106;
RA de Groot N.G., Otting N., Arguello R., Watkins D.I., Doxiadis G.G.M.,
RA Madrigal J.A., Bontrou R.E.;
RT "Major histocompatibility complex class I diversity in a West African
RT chimpanzee population: implications for HIV research.";
RL Immunogenetics 51:398-409(2000).
CC -I- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -I- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
CC EMBL; AF115459; AAF02438.1; -.
DR HSSP; Q95352; 1HHK.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_c1.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; MHC; Transmembrane.
SQ SEQUENCE 365 AA; 40819 MW; 4E95F08E33479E38 CRC64;
Query Match 33.9%; Score 514; DB 7; Length 365;
Best Local Similarity 40.1%; Pred. No. 6.5e-40;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;
QY 5 SHSLHFLPMGASQDGLSLFEALGYVDOLFFVFDHE--SRVPRTPWVSRRSSQW 62
Db 26 SHSMRYFTSVSRPGRGEPRFTAVGYVDDTQFVRFDSDAASQRMPEPRAPWIEQ-GPEYW 84
QY 63 LQLSQSLKGDHMTVDFTWIMENHNASKS-SHTLQVILGCEMQEDNS-TEGYWKYGYDG 120
Db 85 DQETRAKSAHSQTDRLDGLTGRYNGSEAGSHITQIMYGCVDGSDGRFLRGYRDQAYDG 144
QY 121 QDALEFCFDDTLDRAAEPRAPWPTKLEWERHKIRARQNAYLERDCPAQLQQLLELGRGVLD 180

Db 145 KDYIALNEDLRSWTAADMAAQITKRKWEAAH-AAEQORAYLEGTCEWLRRLRYLENGKETL 203
Qy 181 DQOVPLVKVTHH-VTSSVTLRCRALNYYPQNTWKWLKDQKQMDAKEPEPKDVLPNGD 239
Db 204 QRTDPPKTHHHPISDHEATLRCWALGFYPAEITLTWQRDGED-QTQDTLVELVETRPAGD 262
Qy 240 GTYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
Db 263 GTFQKWAAVVVPSEGEQRYTCHVQHEGLPKPLTLRWE 299

Search completed: August 5, 2003, 13:10:01
Job time : 33 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:05:29 ; Search time 38 Seconds
(without alignments)
1152.856 Million cell updates/sec

Title: US-10-092-404-3

Perfect score: 1514

Sequence: 1 RLLRSHLHFLMGASEQDL.....RYTCQVHEPGLDPLIVWE 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1514	100.0	276	20 AAW94297	HFE mutant (H111A/
2	1514	100.0	276	24 AEG72687	Human haemochromat
3	1502	99.2	276	20 AAW94295	Wild-type HFE poly
4	1502	99.2	276	24 AEG72685	Human haemochromat
5	1502	99.2	348	18 AAW36499	Hereditary haemoch
6	1502	99.2	348	21 AAB19149	A human histocoma
7	1502	99.2	348	22 AAB36869	Human hereditary h
8	1497	98.9	438	23 AAU80035	Beta 2 microglobul
9	1493	98.6	276	20 AAW94296	HFE mutant (H63D-H

10	1493	98.6	276	24 AEG72686	Human haemochromat
11	1493	98.6	348	22 AAB36871	Human hereditary h
12	1491	98.5	348	22 AAB36870	Human hereditary h
13	1482	97.9	348	22 AAB36872	Human hereditary h
14	517	34.1	361	22 AAB36873	Rabbit leukocyte a
15	508	33.6	365	22 AAB36874	MHC class I protei
16	504	33.3	92	24 ABP68379	Human colon specif
17	500	33.0	274	21 AAY68275	Human leukocyte an
18	500	33.0	274	21 AAY52929	Human leukocyte an
19	500	33.0	274	22 AAB58690	HLA-A2/A28 family
20	500	33.0	280	22 AAB58690	HLA-A2/A28 protein
21	500	33.0	280	22 AAU10225	Human leukocyte an
22	500	33.0	280	24 ABU08672	Human histocompat
23	500	33.0	415	22 AAU10224	Human partial beta
24	499	33.0	365	21 ABU08671	Human single chain
25	499	33.0	365	21 AAY68265	Human leukocyte an
26	499	33.0	365	21 AAY52919	HLA-A2/A28 family
27	499	33.0	365	22 AAB58680	HLA-A2/A28 protein
28	498	32.9	368	22 AAM24017	Human EST encoded
29	498	32.9	274	21 AAY68276	Human leukocyte an
30	498	32.9	274	21 AAY52930	HLA-A2/A28 family
31	498	32.9	365	22 AAB58691	HLA-A2/A28 protein
32	498	32.9	365	21 AAY68268	Human leukocyte an
33	498	32.9	365	21 AAY52922	HLA-A2/A28 family
34	497	32.8	365	22 AAB58683	HLA-A2/A28 protein
35	497	32.8	274	9 AAF80911	Consensus sequence
36	497	32.8	365	21 AAY68267	Human leukocyte an
37	497	32.8	365	21 AAY52921	HLA-A2/A28 family
38	496	32.8	365	22 AAB58682	HLA-A2/A28 protein
39	496	32.8	274	21 AAY68274	Human leukocyte an
40	496	32.8	274	21 AAY52928	HLA-A2/A28 family
41	496	32.8	365	22 AAB58689	HLA-A2/A28 protein
42	496	32.8	365	21 AAY68266	Human leukocyte an
43	496	32.8	365	21 AAY52920	HLA-A2/A28 family
44	495	32.7	365	22 AAB58681	HLA-A2/A28 protein
45	494	32.6	412	19 AAW68385	Chimeric HLA-A2.1/
			274	21 AAY68273	Human leukocyte an

ALIGNMENTS

RESULT 1

AAW94297
ID AAW94297 standard; peptide; 276 AA.

XX AC AAW94297;

XX DT 27-APR-1999 (first entry)

XX DE HFE mutant (H111A/H145A-HFE) polypeptide sequence.

XX DE HFE; beta-2-microglobulin; beta2m; iron overload; hemochromatosis;

KW transfusion; protein replacement therapy; hereditary hemochromatosis;

KW transferrin receptor; iron deficiency; anemia; mutant.

XX OS Synthetic.

XX FH Key

FT Key Location/Qualifiers

FT Misc-difference 2

FT Misc-difference 99

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

FT Misc-difference 123

PF 12-JUN-1998; 98WO-US12436.
 XX
 PR 13-JUN-1997; 97US-0876010.
 XX
 PA (CALY) CALIFORNIA INST OF TECHNOLOGY.
 PA (PROG-) PROGENITOR INC.
 XX
 PI Bjorkman PJ, Feder JN, Schatzman RC;
 XX WPI; 1999-080886/07.
 DR
 XX New treatment of an iron overload disease - comprises use of HFE
 PT polypeptides provided in a complex with full length, wild type human
 PT (2m), useful in protein replacement therapy
 XX
 PS Claim 5; Page 15; 36pp; English.
 XX
 CC The present sequence represents a H111A/H145A-HFE mutant polypeptide.
 CC The HFE polypeptides (AAW94295-297) provided in a complex with full
 CC length, wild type human beta-2-microglobulin (beta2m) form compositions
 CC in the treatment of primary iron overload diseases (e.g.,
 CC hemochromatosis), or other iron overload conditions resulting from
 CC secondary causes (e.g., repeated transfusions). Data regarding the
 CC structure and function correlations of HFE polypeptides is useful in
 CC designing drugs that modulate the HFE gene and HFE activity. The
 CC polypeptides are also useful in protein replacement therapy for
 CC individuals possessing a defective HFE gene (e.g., Hereditary
 CC hemochromatosis). (Antagonists of the polypeptides are also useful in
 CC treating primary and secondary iron overload diseases. The modulators of
 CC the transferrin receptor are useful in treating iron deficiency
 CC conditions such as anemia, and in modulating the amount of iron
 CC transported into a cell. The HFE polypeptides provide a molecular basis
 CC for the relationship between HFE and iron metabolism, which enables
 CC treatment of iron overload and deficiency diseases.
 XX
 SQ Sequence 276 AA;
 Query Match 100.0%; Score 1514; DB 20; Length 276;
 Best Local Similarity 100.0%; Pred. No. 1.1e-132;
 Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
 Db 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
 QY 61 MWLQLSQSLKGDWHMFTVDFTWMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120
 Db 61 MWLQLSQSLKGDWHMFTVDFTWMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120
 QY 121 QDALEFCPDTLDWRAAEPRAWPTKLEWERHKIRARQNRAVLERDCPAQLQQLLELGRGVL 180
 Db 121 QDALEFCPDTLDWRAAEPRAWPTKLEWERHKIRARQNRAVLERDCPAQLQQLLELGRGVL 180
 QY 181 DQOVPLVKVTHHTVSSVTLTLCRALNYPQNITMKWKDKQPMDAKEPEPKDVLPGD 240
 Db 181 DQOVPLVKVTHHTVSSVTLTLCRALNYPQNITMKWKDKQPMDAKEPEPKDVLPGD 240
 QY 241 TYCGWITLAVPPGEQRQYTCQVEHPGLDQLIWIWE 276
 Db 241 TYCGWITLAVPPGEQRQYTCQVEHPGLDQLIWIWE 276
 RESULT 2
 ABG72687
 ID ABG72687 standard; protein; 276 AA.
 XX
 AC ABG72687;
 XX
 DT 05-MAR-2003 (first entry)
 TX
 DE Human haemochromatosis (HFE) mature protein, mutant H89A/H123A.
 DE
 XX Human; haemochromatosis; HFE; hereditary haemochromatosis;
 KW

KW iron overload disease; iron deficiency disease; Beta2-microglobulin;
 KW Beta2m; transferrin receptor; anaemia; mutant; mutein.
 XX
 OS Homo sapiens.
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT Misc-difference 89 /note= "Wild-type His substituted by Ala"
 FT Misc-difference 123 /note= "Wild-type His substituted by Ala"
 FT
 XX US6391852-B1.
 XX 21-MAY-2002.
 XX
 XX 12-JUN-1998; 98US-0094964.
 XX
 PR 13-JUN-1997; 97US-0876010.
 XX
 PA (BIRA) BIO-RAD LAB INC.
 PA (CALY) CALIFORNIA INST OF TECHNOLOGY.
 XX
 PI Feder JN, Bjorkman PJ, Schatzman RC;
 XX WPI; 2003-155377/15.
 XX
 CC Method of treating an iron overload disease comprises administration of
 CC a soluble complex comprising a 276 amino acid HFE polypeptide and a
 CC full length, wild-type human beta2m -
 CC Claim 3; Column 2; 17pp; English.
 XX
 CC The invention relates to a method of treating an iron overload disease
 CC comprising administration of a soluble complex comprising a 276 amino
 CC acid mature HFE (hereditary haemochromatosis gene protein) polypeptide
 CC (ABG72687) and a full length, wild-type human beta2m
 CC (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of
 CC 1,2⁵¹I-transferrin in the presence of purified H63D-HFE/beta2m
 CC heterodimers was determined. At a concentration of 250 nM H63D-HFE/
 CC beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for
 CC transferrin of 28 nM. At the same concentration of normal HFE/beta 2m
 CC heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence
 CC of any HFE/beta 2m heterodimers, TfR displayed a KD for transferrin of
 CC 7nM. It was observed that H63D-HFE/beta 2m heterodimers were 30-40 % less
 CC efficient in decreasing TfR affinity for transferrin compared to
 CC wild-type HFE. The method is useful for treating iron overload diseases
 CC and iron deficiency e.g. anaemia. The present sequence is the H11A/H145A
 CC (residues 111 and 145 of the full length protein, 89/123 of the mature
 CC form) mutant from of mature HFE used to investigate the role of the His
 CC residues in transferrin receptor binding to transferrin.
 XX
 SQ Sequence 276 AA;
 Query Match 100.0%; Score 1514; DB 24; Length 276;
 Best Local Similarity 100.0%; Pred. No. 1.1e-132;
 Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
 Db 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
 QY 61 MWLQLSQSLKGDWHMFTVDFTWMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120
 Db 61 MWLQLSQSLKGDWHMFTVDFTWMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120
 QY 121 QDALEFCPDTLDWRAAEPRAWPTKLEWERHKIRARQNRAVLERDCPAQLQQLLELGRGVL 180
 Db 121 QDALEFCPDTLDWRAAEPRAWPTKLEWERHKIRARQNRAVLERDCPAQLQQLLELGRGVL 180
 QY 181 DQOVPLVKVTHHTVSSVTLTLCRALNYPQNITMKWKDKQPMDAKEPEPKDVLPGD 240
 Db 181 DQOVPLVKVTHHTVSSVTLTLCRALNYPQNITMKWKDKQPMDAKEPEPKDVLPGD 240

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QY      241 TYQGWTTLAVPPGEGORYTCQVEHPGLDQPLIVWE 276
Db      241 TYQGWTTLAVPPGEGORYTCQVEHPGLDQPLIVWE 276

RESULT 3
AAW94295
ID      AAW94295 standard; peptide; 276 AA.
XX      AAW94295;
XX      27-APR-1999 (first entry)
XX      Wild-type HFE polypeptide sequence.
DE      HFE; beta-2-microglobulin; beta2m; iron overload; hemochromatosis;
KW      transfusion; protein replacement therapy; hereditary hemochromatosis;
KW      transferrin receptor; iron deficiency; anemia.
XX      Unidentified.
OS      XX
FH      Key Location/Qualifiers
FT      Misc-difference 2 /note= "indicated in the sequence listing as Arg"
PT      XX
XX      W09856814-A1.
XX      17-DEC-1998.
XX      12-JUN-1998; 98WO-US12436.
XX      13-JUN-1997; 97US-0876010.
XX      (CALY ) CALIFORNIA INST OF TECHNOLOGY.
PA      (PROG-) PROGENITOR INC.
XX      Bjorkman PJ, Feder JN, Schatzman RC;
PI      WPI; 1999-080886/07.
XX      New treatment of an iron overload disease - comprises use of HFE
PT      polypeptides provided in a complex with full length, wild type human
PT      (2m), useful in protein replacement therapy
XX      Claim 1; Page 13; 36pp; English.
XX      The present sequence represents a wild-type HFE polypeptide. The HFE
CC      polypeptides (AAW94295-297) provided in a complex with full length,
CC      wild type human beta-2-microglobulin (beta2m) form compositions in the
CC      treatment of primary iron overload diseases (e.g. hemochromatosis), or
CC      other iron overload conditions resulting from secondary causes (e.g.
CC      repeated transfusions). Data regarding the structure and function
CC      correlations of HFE polypeptides is useful in designing drugs that
CC      modulate the HFE gene and HFE activity. The polypeptides are also useful
CC      in protein replacement therapy for individuals possessing a defective
CC      HFE gene (e.g. Hereditary hemochromatosis). (Ant)agonists of the
CC      polypeptides are also useful in treating primary and secondary iron
CC      overload diseases. The modulators of the transferrin receptor are useful
CC      in treating iron deficiency conditions such as anemia, and in modulating
CC      the amount of iron transported into a cell. The HFE polypeptides provide
CC      a molecular basis for the relationship between HFE and iron metabolism,
CC      which enables treatment of iron overload and deficiency diseases.
XX      Sequence 276 AA;
SQ      Query Match 99.2%; Score 1502; DB 20; Length 276;
        Best Local Similarity 99.3%; Pred. No. 1.5e-131;
        Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY      1 RLLRSHLYLFWGASEQDLGLSLFALGVDDQLFVFDHESRRVPRTPWVSSRISQ 60
Db      1 RLLRSHLYLFWGASEQDLGLSLFALGVDDQLFVFDHESRRVPRTPWVSSRISQ 60

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QY      61 MMLQLSGLKGDHMFVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGDG 120
Db      61 MMLQLSGLKGDHMFVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGDG 120

QY      121 QDALFECPDTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLDRDCPAQLQQLLELGRGVL 180
Db      121 QDHLSECPDTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLDRDCPAQLQQLLELGRGVL 180

QY      181 DQOVPLVKVTHVHTSSVTTLCRALNYYYPQNTMKWLKDKQPMDAKEPEPKDVLPGDGG 240
Db      181 DQOVPLVKVTHVHTSSVTTLCRALNYYYPQNTMKWLKDKQPMDAKEPEPKDVLPGDGG 240

QY      241 TYQGWTTLAVPPGEGORYTCQVEHPGLDQPLIVWE 276
Db      241 TYQGWTTLAVPPGEGORYTCQVEHPGLDQPLIVWE 276

RESULT 4
ABG72685
ID      ABG72685 standard; protein; 276 AA.
XX      AC ABG72685;
XX      05-MAR-2003 (first entry)
XX      Human haemochromatosis (HFE) mature protein.
XX      Human; haemochromatosis; HFE; hereditary haemochromatosis;
KW      iron overload disease; iron deficiency disease; Beta2-microglobulin;
KW      Beta2m; transferrin receptor; anaemia.
XX      Homo sapiens.
XX      US6391852-B1.
XX      21-MAY-2002.
XX      12-JUN-1998; 98US-0094964.
XX      13-JUN-1997; 97US-0876010.
XX      (BIRA ) BIO-RAD LAB INC.
PA      (CALY ) CALIFORNIA INST OF TECHNOLOGY.
XX      Feder JN, Bjorkman PJ, Schatzman RC;
XX      WPI; 2003-155377/15.
XX      Method of treating an iron overload disease comprises administration of
PT      a soluble complex comprising a 276 amino acid HFE polypeptide and a
PT      full length, wild-type human beta2m -
XX      Claim 1; Column 1; 17pp; English.
XX      The invention relates to a method of treating an iron overload disease
CC      comprising administration of a soluble complex comprising a 276 amino
CC      acid mature HFE (hereditary haemochromatosis gene protein) polypeptide
CC      (ABG72685-ABG72687) and a full length, wild-type human beta2m
CC      (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of
CC      125I-transferrin in the presence of purified H63D-HFE/beta2m
CC      heterodimers was determined. At a concentration of 250 nM H63D-HFE/
CC      beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for
CC      transferrin of 28 nM. At the same concentration of normal HFE/beta 2m
CC      heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence
CC      of any HFE/beta2m heterodimers, TfR displayed a KD for transferrin of
CC      7nM. It was observed that H63D-HFE/beta 2m heterodimers were 30-40 % less
CC      efficient in decreasing TfR affinity for transferrin compared to
CC      wild-type HFE. The method is useful for treating iron overload diseases
CC      and iron deficiency e.g. anaemia. The present sequence is wild-type
XX      mature HFE.
XX      Sequence 276 AA;
SQ

```

Query Match 99.2%; Score 1502; DB 24; Length 276;
 Best Local Similarity 99.3%; Pred. No. 1.5e-131;
 Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFVFDHESRRVPEPTPWSSRISSQ 60
 DB 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFVFDHESRRVPEPTPWSSRISSQ 60

QY 61 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMOEDNSTEGWKYGYDG 120
 DB 61 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMOEDNSTEGWKYGYDG 120

QY 121 QDALEFCPDTLDWRAAPRAWPVKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
 DB 121 QDHLEFCPDTLDWRAAPRAWPVKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

QY 181 DQVPPPLVKVTHHTVSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
 DB 181 DQVPPPLVKVTHHTVSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240

QY 241 TYQGWITLAVPPGEORVTCQVEHPGLDQPLIVWE 276
 DB 241 TYQGWITLAVPPGEORVTCQVEHPGLDQPLIVWE 276

RESULT 5

AAW36499
 ID AAW36499 standard; Protein; 348 AA.

AC AAW36499;

DT 14-APR-1998 (first entry)

DE Hereditary haemochromatosis gene product.

XX Hereditary haemochromatosis; metal toxicity; diagnosis;

XX gene therapy; prenatal screening; human.

XX Homo sapiens.

XX Key Location/Qualifiers

FT Misc-difference 63 /note= "substituted by Asp in 24s2 mutant"

FT Misc-difference 65 /note= "substituted by Cys in 24d7 variant"

FT Misc-difference 282 /note= "substituted by Tyr in 24d1 mutant"

FT WO9738137-A1.

PN 16-OCT-1997.

XX 04-APR-1997; 97WO-US06254.

XX 23-MAY-1996; 96US-0652265.

PR 04-APR-1996; 96US-0630912.

PR 16-APR-1996; 96US-0632673.

XX (MERC-) MERCATOR GENETICS INC.

PA Drayna DT, Feder JN, Gnirke A, Ruddy D, Thomas WJ;

PI Tauchinashi Z, Wolff RK;

XX WPI; 1997-512743/47.

DR N-PSDB; AAT96690, AAT96691.

XX Hereditary haemochromatosis gene and variants - useful for diagnosis
 and treatment of hereditary haemochromatosis disease

XX Disclosure; Fig 4; 115pp; English.

XX This polypeptide is the expression product of a novel human gene

CC (see AAT96690) whose mutated form is associated with hereditary
 haemochromatosis (HH). A single mutation (24d1) in the HH gene
 appears responsible for the majority of HH disease. This comprises
 a G to A substitution that is present in 86% of affected
 chromosomes and in 4% of unaffected chromosomes. It results in a
 Cys to Tyr substitution in the encoded protein at a critical
 disulphide bridge important for secondary structure. The following
 are claimed: the 10825 bp genomic DNA sequence (I), a 1437 bp cDNA
 sequence (Ia) (see AAT96691) and their 24d1, 24d2 and 24d7 variants;
 a cloning or expression vector; host cells; a peptide product
 chosen from the HH gene product, its variants (24d1, 24d2 and
 24d7), or a peptide of at least 56 amino acid residues of these; an
 antibody produced using the peptide as an immunogen; a method to
 determine the presence or absence of the common HH gene mutation;
 an animal model for the HH disease; metal chelation agents; T-cell
 differentiation factors and therapeutic agents for the mitigation
 of injury due to oxidative process in vivo or mitigation of iron
 overload; a method for screening potential therapeutic agents for
 activity in connection with HH disease; an antisense oligonucleotide
 directed against a transcriptional product of a nucleic acid
 sequence as above; and oligonucleotides or pairs of oligonucleotides
 covering a range of nucleotides from (I), (Ia) or their variants;
 useful for detecting a polymorphism in the HH gene. The invention
 also relates to methods for screening for HH homozygotes, to HH
 diagnosis, prenatal screening and diagnosis, and therapies of HH
 disease, including gene therapy, protein- and antibody-based
 therapeutics, and small molecule therapeutics.

XX Sequence 348 AA;

Query Match 99.2%; Score 1502; DB 18; Length 348;

Best Local Similarity 99.3%; Pred. No. 2e-131;

Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFVFDHESRRVPEPTPWSSRISSQ 60

DB 23 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFVFDHESRRVPEPTPWSSRISSQ 82

QY 61 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMOEDNSTEGWKYGYDG 120

DB 83 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMOEDNSTEGWKYGYDG 142

QY 121 QDALEFCPDTLDWRAAPRAWPVKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

DB 143 QDHLEFCPDTLDWRAAPRAWPVKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202

QY 181 DQVPPPLVKVTHHTVSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240

DB 203 DQVPPPLVKVTHHTVSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 262

QY 241 TYQGWITLAVPPGEORVTCQVEHPGLDQPLIVWE 276

DB 263 TYQGWITLAVPPGEORVTCQVEHPGLDQPLIVWE 298

RESULT 6

AAW19149

ID AAW19149 standard; Protein; 348 AA.

XX AAW19149;

DT 19-FEB-2001 (first entry)

DE A human histocompatibility iron loading (HFE) protein.

XX Human; histocompatibility iron loading protein; HFE protein;
 major histocompatibility complex; non-classical class I gene;
 chromosome 6p; iron disorder; haemochromatosis.

OS Homo sapiens.

XX Key Location/Qualifiers

FT Peptide 1..22

FT Misc-difference 63 /note= "signal peptide"
 FT "when nucleotide 187 is mutated to G, then
 FT this residue is Asp"
 FT Misc-difference 65 /note= "when nucleotide 193 is mutated to T, then
 FT this residue is Cys"
 FT Domain 80..108
 FT /note= "alpha domain"
 FT Misc-difference 93 /note= "when nucleotide 277 is mutated to C, then
 FT this residue is Arg"
 FT Misc-difference 105 /note= "when nucleotide 314 is mutated to C, then
 FT this residue is Thr"
 FT
 XX WO200058515-A1.
 XX
 XX 05-OCT-2000.
 XX
 XX 24-MAR-2000; 2000WO-US07982.
 XX
 XX 26-MAR-1999; 99US-0277457.
 XX
 XX (BILL-) BILLUPS-ROTHENBERG INC.
 XX
 XX Rothenberg BE, Sawada-Hirai R, Barton JC;
 XX
 XX WPI; 2000-647244/62.
 XX N-PSDB; AAA96769.
 XX
 XX Diagnosing an iron disorder e.g. hemochromatosis or a genetic
 XX susceptibility to develop it, by determining the presence of a mutation
 XX in exon 2 or an intron of a histocompatibility iron loading nucleic
 XX acid -
 XX
 XX Disclosure; Page 3; 55pp; English.
 XX
 XX The present sequence represents a human histocompatibility iron loading
 XX (HFE) protein. The HFE gene is a major histocompatibility (MHC)
 XX non-classical class I gene located on chromosome 6p. Mutations in the
 XX gene lead to iron disorders. The specification describes a method for
 XX diagnosing an iron disorder or a genetic susceptibility to develop the
 XX disorder in a mammal. The method comprises determining the presence of
 XX a mutation in exon 2 or an intron of a HFE gene or protein. The mutation
 XX is not a C to G missense mutation at nucleotide 187 of the sequence
 XX given in A96769 (Genbank Accession number U60319). The presence of the
 XX mutation indicates the disorder or the genetic susceptibility to the
 XX disorder. The method is used to diagnose an iron disorder
 XX e.g. haemochromatosis, or a genetic susceptibility to develop it.
 XX
 XX Sequence 348 AA;
 SQ
 Query Match 99.2%; Score 1502; DB 21; Length 348;
 Best Local Similarity 99.3%; Pred. No. 2e-131;
 Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 RLRLSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
 DB 23 RLRLSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 82
 QY 61 MWLQLSLSKLGWDHMTVDFTWIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120
 DB 83 MWLQLSLSKLGWDHMTVDFTWIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 142
 QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 180
 DB 143 QDHLEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 202
 QY 181 DQOVPLVKVTHHTVSSVTLRLCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
 DB 203 DQOVPLVKVTHHTVSSVTLRLCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262

QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
 DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298
 RESULT 7
 ID AAB36869
 XX AAB36869 standard; Protein; 348 AA.
 AC AAB36869;
 XX 21-FEB-2001 (first entry)
 DT
 XX Human hereditary hemochromatosis protein.
 DE
 XX HH; hereditary hemochromatosis; chelation agent;
 KW T-cell differentiation factor; iron overload.
 XX
 XX Homo sapiens.
 OS
 XX US6140305-A.
 PN
 XX 31-OCT-2000.
 PD
 XX 04-APR-1997; 97US-0834497.
 XX
 XX 04-APR-1996; 96US-0630912.
 PR 16-APR-1996; 96US-0632673.
 PR 23-MAY-1996; 96US-0652265.
 XX
 XX (BIRA) BIO-RAD LAB INC.
 PA
 XX Thomas WJ, Drayna DT, Gnikre A, Ruddy D, Tsuchihashi Z, Wolff RK;
 PI Feder JN;
 XX
 XX WPI; 2001-006341/01.
 DR N-PSDB; AAC68425.
 XX
 XX New hereditary hemochromatosis gene products or polypeptides, useful
 XX for treating hereditary hemochromatosis in a patient, and as a metal
 XX chelation agent alleviating iron overload -
 XX
 XX Claim 1; Fig 4; 108pp; English.
 XX
 XX The present invention relates to hereditary hemochromatosis gene
 XX products. These proteins may be used to treat a patient diagnosed as
 XX having human hemochromatosis disease. It is also useful as a metal
 XX chelation agent or as a T-cell differentiation factor, and for
 XX alleviating iron overload. They may also be used in protein replacement
 XX therapy for individuals having a defective human hemochromatosis gene.
 XX
 XX Sequence 348 AA;
 SQ
 Query Match 99.2%; Score 1502; DB 22; Length 348;
 Best Local Similarity 99.3%; Pred. No. 2e-131;
 Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 1 RLRLSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
 DB 23 RLRLSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 82
 QY 61 MWLQLSLSKLGWDHMTVDFTWIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120
 DB 83 MWLQLSLSKLGWDHMTVDFTWIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 142
 QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 180
 DB 143 QDHLEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 202
 QY 181 DQOVPLVKVTHHTVSSVTLRLCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
 DB 203 DQOVPLVKVTHHTVSSVTLRLCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262

QY 241 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIVWE 276
 |||||
 Db 263 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIVWE 298
 |||||

RESULT 8

AAU80035
 ID AAU80035 standard; Protein; 438 AA.

AC AAU80035;

DT 15-JUL-2002 (first entry)

DE Beta 2 microglobulin (beta2m)/HFE monochain.

XX Human; beta 2 microglobulin; beta2m/HFE monochain; HFE; ischaemia;
 KW iron absorption regulator; intracellular iron absorption; lung injury;
 KW haemochromatosis; transfusion; thalassaemia; haemolytic anaemia;
 KW chronic infection; transferrin receptor; Tfr; brain tumour; cancer;
 KW oxidative stress disorder; tissue damage; vascular disease;
 KW inflammation; atherosclerosis; autoimmune disease;
 KW inflammatory condition.

XX Homo sapiens.

XX WO200224929-A2.

XX 28-MAR-2002.

XX 24-SEP-2001; 2001WO-US29873.

XX 22-SEP-2000; 2000US-234843P.

XX (UYRA-) UNIV RAMOT APPLIED RES & IND DEV LTD.
 PA (MCIN/) MCINNIS P.

PI Ehrlich R, Rotem-Yehudar R, Laham N;

XX WPI; 2002-383192/41.

DR N-PSDB; ABK49917.

XX Soluble beta 2 microglobulin/HFE monochain useful for treating
 PT iron-overload conditions e.g. thalassaemia and chronic infections,
 PT comprises human beta 2 microglobulin linked to alpha domains of HFE by
 PT a linker peptide

XX Example 2; Fig 2; 77pp; English.

XX The invention relates to a soluble polypeptide (I) of beta 2
 CC microglobulin (beta2m)/HFE monochain comprising human beta2m (or its
 CC analogue or active fragment), linked to alpha1-alpha3 domains of human
 CC HFE (a central regulator of iron absorption; undefined), or its analogue
 CC or active fragment, by a flexible linker peptide, or a functional
 CC derivative or salt of (I). (I) is useful for reducing intracellular iron
 CC absorption in patients having hereditary haemochromatosis, transfusions,
 CC thalassaemias, haemolytic anaemia or chronic infections, and for
 CC delivering a therapeutic to cells that over-express transferrin receptor
 CC (Tfr) which are preferably lymphocytes or leukocytes, across the blood-
 CC brain barrier. (I) is further useful for treating brain tumour. (I)
 CC is also useful for treating oxidative stress disorders resulting in
 CC tissue damage e.g. vascular diseases, inflammation, atherosclerosis,
 CC lung injury, ischaemia, etc. A DNA molecule (II) encoding (I) is useful
 CC as a platform for drug delivery of therapeutic use for cancer,
 CC autoimmune diseases and inflammatory conditions. The monochain manifests
 CC specific characteristics advantageous for drug delivery systems. It is a
 CC soluble, stable and fully conformed protein. It binds specifically to
 CC transferrin receptor (Tfr) and therefore targets cells that over-express
 CC this receptor. It is continuously internalised by the target cells, thus
 CC enabling efficient drug delivery. It dissociates from the receptor in the
 CC cells, minimising side effects. It negatively regulates iron absorption,
 CC reducing growth of undesired cells and preventing lymphocyte activation.
 CC It is not diluted in the blood as is transferrin. It should not induce an
 CC immune response since it is a self non-polymorphic protein and delivery of

CC drugs via monochain is expected to overcome drug-resistance since it is a
 CC natural Tfr-binding protein. The present sequence represents the amino
 CC acid sequence of beta2m/HFE monochain.

XX SQ Sequence 438 AA;

Query Match 98.9%; Score 1497; DB 23; Length 438;

Best Local Similarity 99.3%; Pred. No. 7.8e-131;

Matches 273; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYDDQLFVFDHESRRRVEPRTPWSSRISSQ 60
 |||||

Db 135 RLLRSHSLHYLFMGASEQDLGLSLFEALGYDDQLFVFDHESRRRVEPRTPWSSRISSQ 194
 |||||

QY 61 MWLQLSQSLKGDWHMFTVDFWTIMENHNASKESHTLQVILGCEMEDNSTEGYWKYGYDG 120
 |||||

Db 195 MWLQLSQSLKGDWHMFTVDFWTIMENHNASKESHTLQVILGCEMEDNSTEGYWKYGYDG 254
 |||||

QY 121 QDALEFCPDTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
 |||||

Db 255 QDHLSEFCPDTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 314
 |||||

QY 181 DQVPLVKVTHVTSSVTTLRCRALNYPQNTWKWLKDKQPMDAKEPEPKDVLPGDGG 240
 |||||

Db 315 DQVPLVKVTHVTSSVTTLRCRALNYPQNTWKWLKDKQPMDAKEPEPKDVLPGDGG 374
 |||||

QY 241 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIVW 275
 |||||

Db 375 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIVW 409
 |||||

RESULT 9

AAW94296

ID AAW94296 standard; peptide; 276 AA.

XX AC AAW94296;

XX 27-APR-1999 (first entry)

XX HFE mutant (H63D-HFE) polypeptide sequence.

XX HFE; beta-2-microglobulin; beta2m; iron overload; hemochromatosis;

KW transfusion; protein replacement therapy; hereditary hemochromatosis;

KW transferrin receptor; iron deficiency; anemia; mutant.

XX Synthetic.

XX Key Location/Qualifiers

FT Misc-difference 2

FT /note= "indicated in the sequence listing as Arg"

FT Misc-difference 41

FT /label= H63D

FT /note= "wild type His (of the mature protein sequence)

FT is replaced by Asp"

PN WO9856814-A1.

XX 17-DEC-1998.

XX 12-JUN-1998; 98WO-US12436.

XX 13-JUN-1997; 97US-0876010.

XX (CALY) CALIFORNIA INST OF TECHNOLOGY.

PA (PROG-) PROGENITOR INC.

XX Bjorkman PU, Feder JN, Schatzman RC;

XX WPI; 1999-080886/07.

XX New treatment of an iron overload disease - comprises use of HFE

PT polypeptides provided in a complex with full length, wild type human

PT (2m), useful in protein replacement therapy

XX PS Claim 3; Page 14; 36pp; English.

XX CC The present sequence represents a H63D-HFE mutant polypeptide. The HFE

XX CC polypeptides (AAW94295-297) provided in a complex with full length,

XX CC wild type human beta-2-microglobulin (beta2m) form compositions in the

XX CC treatment of primary iron overload diseases (e.g. hemochromatosis), or

XX CC other iron overload conditions resulting from secondary causes (e.g.

XX CC repeated transfusions). Data regarding the structure and function

XX CC correlate of HFE polypeptides is useful in designing drugs that

XX CC modulate the HFE gene and HFE activity. The polypeptides are also useful

XX CC in protein replacement therapy for individuals possessing a defective

XX CC HFE gene (e.g. Hereditary hemochromatosis). (Antagonists of the

XX CC polypeptides are also useful in treating primary and secondary iron

XX CC overload diseases. The modulators of the transferrin receptor are useful

XX CC in treating iron deficiency conditions such as anemia, and in modulating

XX CC the amount of iron transported into a cell. The HFE polypeptides provide

XX CC a molecular basis for the relationship between HFE and iron metabolism,

XX CC which enables treatment of iron overload and deficiency diseases.

XX SQ Sequence 276 AA;

Query Match 98.6%; Score 1493; DB 20; Length 276;

Best Local Similarity 98.9%; Pred. No. 1e-130;

Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60

DB 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60

QY 61 MWLQSLQSLKGWDMFTVDFWTIMENHNASKESHTLQVLGCEMOEDNSTEGYWKYGYDG 120

DB 61 MWLQSLQSLKGWDMFTVDFWTIMENHNASKESHTLQVLGCEMOEDNSTEGYWKYGYDG 120

QY 121 QDALEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

DB 121 QDHLEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

QY 181 DQOVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEFEKPKDVLPGDVG 240

DB 181 DQOVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEFEKPKDVLPGDVG 240

QY 241 TYQGWITLAVPPGGEQRYTCQVEHPGLDQPLIWIWE 276

DB 241 TYQGWITLAVPPGGEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 10

ABG72686

ID ABG72686 standard; protein; 276 AA.

XX AC ABG72686;

XX DT 05-MAR-2003 (first entry)

XX DE Human haemochromatosis (HFE) mature protein, mutant H41D.

XX KW Human; haemochromatosis; HFE; hereditary haemochromatosis;

XX KW iron overload disease; iron deficiency disease; Beta2-microglobulin;

XX KW Beta2m; transferrin receptor; anaemia; mutant; mutein.

XX OS Homo sapiens.

XX OS Synthetic.

XX FH Key

XX FT Location/Qualifiers

XX FT Misc-difference 41

XX FT /note= "Wild-type His substituted by Asp"

XX PN US6391852-B1.

XX XX 21-MAY-2002.

XX PD 12-JUN-1998; 98US-0094964.

XX PF

XX PR 13-JUN-1997; 97US-0876010.

XX PA (BIRA) BIO-RAD LAB INC.

XX PA (CALY) CALIFORNIA INST OF TECHNOLOGY.

XX PI Feder JN, Bjorkman PJ, Schatzman RC;

XX XX WPI; 2003-155377/15.

XX XX Method of treating an iron overload disease comprises administration of

XX PT a soluble complex comprising a 276 amino acid HFE polypeptide and a

XX PT full length, wild-type human beta2m -

XX PS Claim 2; Column 2; 17pp; English.

XX CC The invention relates to a method of treating an iron overload disease

XX CC comprising administration of a soluble complex comprising a 276 amino

XX CC acid mature HFE (hereditary haemochromatosis gene protein) polypeptide

XX CC (ABG72685-ABG72687) and a full length, wild-type human beta2m

XX CC (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of

XX CC ⁵¹Fe-transferrin in the presence of purified H63D-HFE/beta2m

XX CC heterodimers was determined. At a concentration of 250 nM H63D-HFE/

XX CC beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for

XX CC transferrin of 28 nM. At the same concentration of normal HFE/beta 2m

XX CC heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence

XX CC of any HFE/beta 2m heterodimers, TfR displayed a KD for transferrin of

XX CC 7nM. It was observed that H63D-HFE/beta 2m heterodimers were 30-40 % less

XX CC efficient in decreasing TfR affinity for transferrin compared to

XX CC wild-type HFE. The method is useful for treating iron overload diseases

XX CC and iron deficiency e.g. anaemia. The present sequence is the H63D

XX CC (residue 63 of the full length protein, 41 of the mature form)

XX CC mutant form of mature HFE used to investigate the role of the His

XX CC residue in transferrin receptor binding to transferrin.

XX SQ Sequence 276 AA;

Query Match 98.6%; Score 1493; DB 24; Length 276;

Best Local Similarity 98.9%; Pred. No. 1e-130;

Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60

DB 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60

QY 61 MWLQSLQSLKGWDMFTVDFWTIMENHNASKESHTLQVLGCEMOEDNSTEGYWKYGYDG 120

DB 61 MWLQSLQSLKGWDMFTVDFWTIMENHNASKESHTLQVLGCEMOEDNSTEGYWKYGYDG 120

QY 121 QDALEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

DB 121 QDHLEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

QY 181 DQOVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEFEKPKDVLPGDVG 240

DB 181 DQOVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEFEKPKDVLPGDVG 240

QY 241 TYQGWITLAVPPGGEQRYTCQVEHPGLDQPLIWIWE 276

DB 241 TYQGWITLAVPPGGEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 11

AB36871

ID AAB36871 standard; Protein; 348 AA.

XX AC AAB36871;

XX XX 21-FEB-2001 (first entry)

XX DE Human hereditary hemochromatosis 24d2 mutation protein.

XX KW HH; hereditary hemochromatosis; chelation agent;

KW T-cell differentiation factor; iron overload.

XX Homo sapiens.

OS US6140305-A.

PN 31-OCT-2000.

XX 04-APR-1997; 97US-0834497.

XX 04-APR-1996; 96US-0630912.

PR 16-APR-1996; 96US-0632673.

PR 23-MAY-1996; 96US-0652265.

XX (BIRA) BIO-RAD LAB INC.

XX Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;

PI Feder JN;

XX WPI; 2001-006341/01.

DR N-PSDB; AAC68427.

XX New hereditary hemochromatosis gene products or polypeptides, useful

PT for treating hereditary hemochromatosis in a patient, and as a metal

PT chelation agent alleviating iron overload -

XX Claim 3; Fig 4; 108pp; English.

XX The present invention relates to hereditary hemochromatosis gene

CC products. These proteins may be used to treat a patient diagnosed as

CC having human hemochromatosis disease. It is also useful as a metal

CC chelation agent or as a T-cell differentiation factor, and for

CC alleviating iron overload. They may also be used in protein replacement

CC therapy for individuals having a defective human hemochromatosis gene.

XX SQ Sequence 348 AA;

Query Match 98.6%; Score 1493; DB 22; Length 348;

Best Local Similarity 98.9%; Pred. No. 1.4e-130;

Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVETPTWSSRISSQ 60

DB 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVETPTWSSRISSQ 82

QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNHNSKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120

DB 83 MWLQLSQSLKGDHMTVDFTWIMENHNHNSKESHTLQVILGCEMOEDNSTEGYWKYGYDG 142

QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQRAYLERDQPAQLQELLEGRGVL 180

DB 143 QDHLEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQRAYLERDQPAQLQELLEGRGVL 202

QY 181 DQVPPPLVKVTHVTSVTTLRCAALNYYPQNTMKWLKDKQPMDAKEFEPKDVLPNGDG 240

DB 203 DQVPPPLVKVTHVTSVTTLRCAALNYYPQNTMKWLKDKQPMDAKEFEPKDVLPNGDG 262

QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 12

AAB36870

ID AAB36870 standard; Protein; 348 AA.

XX AAB36870;

XX 21-FEB-2001 (first entry)

XX Human hereditary hemochromatosis 24d1 mutation protein.

XX HH; hereditary hemochromatosis; chelation agent;

KW

KW T-cell differentiation factor; iron overload.

XX Homo sapiens.

OS US6140305-A.

PN 31-OCT-2000.

XX 04-APR-1997; 97US-0834497.

XX 04-APR-1996; 96US-0630912.

PR 16-APR-1996; 96US-0632673.

PR 23-MAY-1996; 96US-0652265.

XX (BIRA) BIO-RAD LAB INC.

XX Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;

PI Feder JN;

XX WPI; 2001-006341/01.

DR N-PSDB; AAC68426.

XX New hereditary hemochromatosis gene products or polypeptides, useful

PT for treating hereditary hemochromatosis in a patient, and as a metal

PT chelation agent alleviating iron overload -

XX Claim 2; Fig 3; 108pp; English.

XX The present invention relates to hereditary hemochromatosis gene

CC products. These proteins may be used to treat a patient diagnosed as

CC having human hemochromatosis disease. It is also useful as a metal

CC chelation agent or as a T-cell differentiation factor, and for

CC alleviating iron overload. They may also be used in protein replacement

CC therapy for individuals having a defective human hemochromatosis gene.

XX SQ Sequence 348 AA;

Query Match 98.5%; Score 1491; DB 22; Length 348;

Best Local Similarity 98.9%; Pred. No. 2.1e-130;

Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVETPTWSSRISSQ 60

DB 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVETPTWSSRISSQ 82

QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNHNSKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120

DB 83 MWLQLSQSLKGDHMTVDFTWIMENHNHNSKESHTLQVILGCEMOEDNSTEGYWKYGYDG 142

QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQRAYLERDQPAQLQELLEGRGVL 180

DB 143 QDHLEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQRAYLERDQPAQLQELLEGRGVL 202

QY 181 DQVPPPLVKVTHVTSVTTLRCAALNYYPQNTMKWLKDKQPMDAKEFEPKDVLPNGDG 240

DB 203 DQVPPPLVKVTHVTSVTTLRCAALNYYPQNTMKWLKDKQPMDAKEFEPKDVLPNGDG 262

QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 13

AAB36872

ID AAB36872 standard; Protein; 348 AA.

XX AAB36872;

XX 21-FEB-2001 (first entry)

XX Human hereditary hemochromatosis 24d1/2 mutation protein.

XX HH; hereditary hemochromatosis; chelation agent;

KW

KW T-cell differentiation factor; iron overload.
 XX Homo sapiens.
 XX US6140305-A.
 XX 31-OCT-2000.
 XX 04-APR-1997; 97US-0834497.
 XX 04-APR-1996; 96US-0630912.
 XX 16-APR-1996; 96US-0632673.
 XX 23-MAY-1996; 96US-0652265.
 XX (BIRA) BIO-RAD LAB INC.
 XX Thomas WJ, Drayna DT, Gnrirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
 PI Feder JN;
 XX WPI; 2001-006341/01.
 XX N-PSDB; AAC68428.
 XX New hereditary hemochromatosis gene products or polypeptides, useful
 PT for treating hereditary hemochromatosis in a patient, and as a metal
 PT chelation agent alleviating iron overload -
 XX
 XX Claim 4; Fig 4; 108pp; English.
 XX The present invention relates to hereditary hemochromatosis gene
 CC products. These proteins may be used to treat a patient diagnosed as
 CC having human hemochromatosis disease. It is also useful as a metal
 CC chelation agent or as a T-cell differentiation factor, and for
 CC alleviating iron overload. They may also be used in protein replacement
 CC therapy for individuals having a defective human hemochromatosis gene.
 XX
 XX Sequence 348 AA;
 SQ
 Query Match 97.9%; Score 1482; DB 22; Length 348;
 Best Local Similarity 98.6%; Pred. No. 1.5e-129;
 Matches 272; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
 QY 1 RLLRSHLYLFGASEQDGLSLFEALGYDDQLFVFDHESRRVPRTPWYSSRISSQ 60
 Db 23 RLLRSHLYLFGASEQDGLSLFEALGYDDQLFVFDHESRRVPRTPWYSSRISSQ 82
 QY 61 MWLQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNS-TEGYWKYGYDG 120
 Db 83 MWLQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNS-TEGYWKYGYDG 142
 QY 121 QDALEFCPDTLDWRAAEPRAPWPTKLEWRHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
 Db 143 QDHLEFCPDTLDWRAAEPRAPWPTKLEWRHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
 QY 181 DQOVPLVKVTHVTSSTVTLRCALNYPQNTMKWLKDKQPMDAKEFEKPKDVLNPGDG 240
 Db 203 DQOVPLVKVTHVTSSTVTLRCALNYPQNTMKWLKDKQPMDAKEFEKPKDVLNPGDG 262
 QY 241 TYQGWITLAVPPGEORVTCQVHPGLDQPLIWIWE 276
 Db 263 TYQGWITLAVPPGEORVTCQVHPGLDQPLIWIWE 298
 RESULT 14
 AAB36873
 ID AAB36873 standard; Protein; 361 AA.
 XX
 AC AAB36873;
 XX
 DT 21-FEB-2001 (first entry)
 XX
 DE Rabbit leukocyte antigen.
 XX
 KW HH; hereditary hemochromatosis; chelation agent;

KW T-cell differentiation factor; iron overload.
 XX Oryctolagus cuniculus.
 XX US6140305-A.
 XX 31-OCT-2000.
 XX 04-APR-1997; 97US-0834497.
 XX 04-APR-1996; 96US-0630912.
 XX 16-APR-1996; 96US-0632673.
 XX 23-MAY-1996; 96US-0652265.
 XX (BIRA) BIO-RAD LAB INC.
 XX Thomas WJ, Drayna DT, Gnrirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
 PI Feder JN;
 XX WPI; 2001-006341/01.
 XX New hereditary hemochromatosis gene products or polypeptides, useful
 PT for treating hereditary hemochromatosis in a patient, and as a metal
 PT chelation agent alleviating iron overload -
 XX
 XX Disclosure; Fig 7; 108pp; English.
 XX The present invention relates to hereditary hemochromatosis gene
 CC products. These proteins may be used to treat a patient diagnosed as
 CC having human hemochromatosis disease. It is also useful as a metal
 CC chelation agent or as a T-cell differentiation factor, and for
 CC alleviating iron overload. They may also be used in protein replacement
 CC therapy for individuals having a defective human hemochromatosis gene.
 XX
 XX Sequence 361 AA;
 SQ
 Query Match 34.1%; Score 517; DB 22; Length 361;
 Best Local Similarity 40.1%; Pred. No. 1.1e-39;
 Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;
 QY 5 SHSLHYLFGASEQDGLSLFEALGYDDQLFVFDHESRRVPRTPWYSSRISSQ 62
 Db 26 SHSMRYFTYSVRPGLGEPRFIIVGVDDTQVRFDSDAASPRMEQRAPEWM-QQVEPEY 84
 QY 63 LQLSLSKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNS-TEGYWKYGYDG 120
 Db 85 DQQTQIAKDTATFRVNLNTALRYNQSAAGSHITQTWFGCEVWADGRFFHGYRQYAYDG 144
 QY 121 QDALEFCPDTLDWRAAEPRAPWPTKLEWRHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
 Db 145 ADVIALNEDLRSWTAADTAQNTQKWEAAG-EAERHRAYLERECVEWLLRYLENGKETL 203
 QY 181 DQOVPLVKVTHVTSSTVTLRCALNYPQNTMKWLKDKQPMDAKEFEKPKDVLNPGDG 239
 Db 204 QRADPPKAVHTVHPASDREATLRWALGFYPAEISLTWQDGED-QTQDTLVELTRPGDG 262
 QY 240 TYQGWITLAVPPGEORVTCQVHPGLDQPLIWIWE 276
 Db 263 GTFOKAAVVPGEORVTCQVHPGLDQPLIWIWE 299
 RESULT 15
 AAB36874
 ID AAB36874 standard; Protein; 365 AA.
 XX
 AC AAB36874;
 XX
 DT 21-FEB-2001 (first entry)
 XX
 DE MHC class I protein.
 XX
 KW HH; hereditary hemochromatosis; chelation agent;
 KW T-cell differentiation factor; iron overload.

[illegible]

Search completed: August 5, 2003, 13:08:24
Job time : 39 secs

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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:10:04 ; Search time 33 Seconds
(without alignments)
993.264 Million cell updates/sec

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Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 451899 seqs, 118759770 residues

Total number of hits satisfying chosen parameters: 451899

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA.*
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8: /cgn2_6/ptodata/1/pubpaa/US08_PUBCOMB.pep.*
9: /cgn2_6/ptodata/1/pubpaa/US09A_PUBCOMB.pep.*
10: /cgn2_6/ptodata/1/pubpaa/US09B_PUBCOMB.pep.*
11: /cgn2_6/ptodata/1/pubpaa/US09C_PUBCOMB.pep.*
12: /cgn2_6/ptodata/1/pubpaa/US09_NEW_PUB.pep.*
13: /cgn2_6/ptodata/1/pubpaa/US10A_PUBCOMB.pep.*
14: /cgn2_6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1514	100.0	276	15	US-10-092-404-3
2	1502	99.2	276	15	US-10-092-404-1
3	1502	99.2	348	12	US-09-981-606-2
4	1493	98.6	276	15	US-10-092-404-2
5	504	33.3	92	14	US-10-016-634A-120
6	500	33.0	280	15	US-10-073-300-6
7	500	33.0	415	15	US-10-073-300-5
8	486	32.1	298	15	US-10-205-823-40
9	486	32.1	298	15	US-10-205-823-42
10	486	32.1	298	15	US-10-177-293-23
11	471	31.1	542	15	US-10-015-535-32
12	471	31.1	542	15	US-10-015-535-34
13	470	31.0	542	15	US-10-015-535-36
14	468	30.9	540	15	US-10-015-535-22
15	468	30.9	541	15	US-10-015-535-28

16	468	30.9	542	15	US-10-015-535-24	Sequence 24, Appl
17	468	30.9	542	15	US-10-015-535-26	Sequence 26, Appl
18	446	29.5	332	9	US-09-870-521-3	Sequence 3, Appl
19	441	29.1	334	9	US-09-870-521-4	Sequence 4, Appl
20	439	29.0	540	15	US-10-015-535-30	Sequence 30, Appl
21	354.5	23.4	170	9	US-09-925-301-1307	Sequence 1307, Appl
22	330	21.8	271	9	US-09-925-301-1431	Sequence 1431, Appl
23	275	18.2	145	9	US-09-810-560-8	Sequence 8, Appl
24	273	18.0	181	11	US-09-013-077A-13	Sequence 13, Appl
25	237	15.7	184	10	US-09-858-580-21	Sequence 21, Appl
26	237	15.7	184	11	US-09-847-172-21	Sequence 21, Appl
27	226	14.9	91	9	US-09-864-761-38005	Sequence 38005, A
28	223	14.7	91	9	US-09-864-761-35461	Sequence 35461, A
29	217.5	14.4	171	15	US-10-144-929-116	Sequence 116, Appl
30	210.5	13.9	104	9	US-09-925-302-835	Sequence 835, Appl
31	207	13.7	117	9	US-09-810-560-9	Sequence 9, Appl
32	202.5	13.4	183	15	US-10-036-542-62	Sequence 62, Appl
33	196.5	13.0	93	9	US-09-864-761-39479	Sequence 39479, A
34	196.5	13.0	110	9	US-09-864-761-35339	Sequence 35339, A
35	196.5	13.0	114	9	US-09-864-761-37988	Sequence 37988, A
36	176.5	11.7	261	10	US-09-925-664-30	Sequence 30, Appl
37	173	11.4	110	10	US-09-796-692-799	Sequence 799, Appl
38	173	11.4	110	10	US-09-796-692-2139	Sequence 2139, Appl
39	173	11.4	110	15	US-10-040-862-799	Sequence 799, Appl
40	173	11.4	110	15	US-10-040-862-2139	Sequence 2139, Appl
41	170	11.2	411	14	US-10-015-536-17	Sequence 17, Appl
42	167.5	11.1	285	10	US-09-756-983-24	Sequence 24, Appl
43	167	11.0	246	9	US-09-989-722-225	Sequence 225, Appl
44	167	11.0	246	9	US-09-989-723-225	Sequence 225, Appl
45	167	11.0	246	9	US-09-989-279-225	Sequence 225, Appl

ALIGNMENTS

RESULT 1

US-10-092-404-3

; Sequence 3, Application US/10092404

; Publication No. US20030073627A1

; GENERAL INFORMATION:

; APPLICANT: Feder, John N.

; Bjorkman, Pamela J.

; Schatzman, Randall C.

; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR

; DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES

; AND IRON DEFICIENCY DISEASES

; NUMBER OF SEQUENCES: 5

; CORRESPONDENCE ADDRESS:

; ADDRESSER: Pennie & Edmonds, LLP

; STREET: 1155 Avenue of the Americas

; CITY: New York

; STATE: NY

; COUNTRY: USA

; ZIP: 10036-2811

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Diskette

; OPERATING SYSTEM: Windows

; SOFTWARE: FastSeq for Windows Version 2.0b

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/10/092,404

; FILING DATE: 04-Mar-2002

; CLASSIFICATION: <Unknown>

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US/09/094,964

; FILING DATE: June 12, 1998

; APPLICATION NUMBER: 08/876,010

; FILING DATE: June 13, 1997

; ATTORNEY/AGENT INFORMATION:

; NAME: Poissant, Brian M

; REGISTRATION NUMBER: 28,462

; REFERENCE/DOCKET NUMBER: 8907-0074-999

; TELECOMMUNICATION INFORMATION:

TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 3:
US-10-092-404-3

Query Match 100.0%; Score 1514; DB 15; Length 276;
Best Local Similarity 100.0%; Pred. No. 1.8e-147;
Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60
QY 61 MWLQSLQSLKGDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQSLQSLKGDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDALEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 121 QDALEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
DB 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 2

US-10-092-404-1
Sequence 1, Application US/10092404
Publication No. US20030073627A1
GENERAL INFORMATION:
APPLICANT: Feder, John N.
Bjorkman, Pamela J.
Schatzman, Randall C.
TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
AND IRON DEFICIENCY DISEASES
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds, LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows
SOFTWARE: FastSeq for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/10/092,404
FILING DATE: 04-Mar-2002
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/09/094,964
FILING DATE: June 12, 1998
APPLICATION NUMBER: 08/876,010
FILING DATE: June 13, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M

REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0074-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-10-092-404-1

Query Match 99.2%; Score 1502; DB 15; Length 276;
Best Local Similarity 99.3%; Pred. No. 3.1e-146;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60
QY 61 MWLQSLQSLKGDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQSLQSLKGDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDALEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 121 QDALEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
DB 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 3

US-09-981-606-2
Sequence 2, Application US/09981606
Publication No. US20030129595A1
GENERAL INFORMATION:
APPLICANT: Rothenberg et al.
TITLE OF INVENTION: Mutations associated with iron disorders
FILE REFERENCE: 24065-004CON
CURRENT APPLICATION NUMBER: US/09/981,606
CURRENT FILING DATE: 2002-10-16
PRIOR APPLICATION NUMBER: 09/277,457
PRIOR FILING DATE: 1999-03-26
NUMBER OF SEQ ID NOS: 30
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 2
LENGTH: 348
TYPE: PRT
ORGANISM: Homo sapiens
US-09-981-606-2

Query Match 99.2%; Score 1502; DB 12; Length 348;
Best Local Similarity 99.3%; Pred. No. 4.3e-146;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 82
QY 61 MWLQSLQSLKGDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142

QY 121 QDALEPCDPTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 180
Db 143 QDLEPCDPTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 240
Db 203 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
Db 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 4
US-10-092-404-2
; Sequence 2, Application US/10092404
; Publication No. US20030073627A1
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; Bjorkman, Pamela J.
; Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; AND IRON DEFICIENCY DISEASES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; OPERATING SYSTEM: Windows
; SOFTWARE: FASTSEQ for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/10/092,404
; FILING DATE: 04-Mar-2002
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 276 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; SEQUENCE DESCRIPTION: SEQ ID NO: 2:
US-10-092-404-2

Query Match 98.6%; Score 1493; DB 15; Length 276;
Best Local Similarity 98.9%; Pred. No. 2.7e-145;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLRLSHSLYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
Db 1 RLRLSHSLYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
QY 61 MMLQLSLSKGDHMTFVDFWTIMENHNASKESHTLQVLGCEMQEDNSTEGYWKYGYDG 120

Db 61 MMLQLSLSKGDHMTFVDFWTIMENHNASKESHTLQVLGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDALEPCDPTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 180
Db 121 QDLEPCDPTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 180
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 240
Db 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
Db 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 5
US-10-016-634A-120
; Sequence 120, Application US/10016634A
; Publication No. US20020192666A1
; GENERAL INFORMATION:
; APPLICANT: Sun, Yongming
; APPLICANT: Recipon, Herve
; APPLICANT: Ghosh, Malavika
; APPLICANT: Liu, Changhai
; TITLE OF INVENTION: Compositions and Methods Relating to Colon Specific Genes and Pri
; FILE REFERENCE: DEX-0255
; CURRENT APPLICATION NUMBER: US/10/016,634A
; CURRENT FILING DATE: 2001-10-31
; PRIOR APPLICATION NUMBER: US 60/244,258
; PRIOR FILING DATE: 2000-10-31
; NUMBER OF SEQ ID NOS: 176
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 120
; LENGTH: 92
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-016-634A-120

Query Match 33.3%; Score 504; DB 14; Length 92;
Best Local Similarity 98.9%; Pred. No. 3.1e-44;
Matches 91; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 92 ESHTLQVLGCEMQEDNSTEGYWKYGYDGQDALEFCPTDLWRAAEPRAPWTKLEWERHK 151
Db 1 ESHTLQVLGCEMQEDNSTEGYWKYGYDGQDHLFCPTDLWRAAEPRAPWTKLEWERHK 60
QY 152 IRARONRAYLERDPCPAQLQQLLELGRGVLDDQ 183
Db 61 IRARONRAYLERDPCPAQLQQLLELGRGVLDDQ 92

RESULT 6
US-10-073-300-6
; Sequence 6, Application US/10073300
; Publication No. US20030003535A1
; GENERAL INFORMATION:
; APPLICANT: Reiter, Yoram
; TITLE OF INVENTION: SINGLE CHAIN CLASS I MAJOR HISTO- COMPATIBILITY COMPLEXES
; FILE REFERENCE: 02/23339
; CURRENT APPLICATION NUMBER: US/10/073,300
; CURRENT FILING DATE: 2002-06-25
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 6
; LENGTH: 280
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-073-300-6

Query Match 33.0%; Score 500; DB 15; Length 280;
Best Local Similarity 39.4%; Pred. No. 3.6e-43;
Matches 109; Conservative 44; Mismatches 116; Indels 8; Gaps 7;

APPLICANT: Zhao, Xumei
TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND
METHODS FOR IDENTIFICATION, ASSESSMENT, PREVENTION, AND
THERAPY OF PROSTATE CANCER
FILE REFERENCE: MRI-044
CURRENT APPLICATION NUMBER: US/10/205,823
CURRENT FILING DATE: 2002-07-25
PRIOR APPLICATION NUMBER: 60/307,982
PRIOR FILING DATE: 2001-07-25
PRIOR APPLICATION NUMBER: 60/314,356
PRIOR FILING DATE: 2001-08-22
PRIOR APPLICATION NUMBER: 60/325,020
PRIOR FILING DATE: 2001-09-25
PRIOR APPLICATION NUMBER: 60/341,746
PRIOR FILING DATE: 2001-12-12
PRIOR APPLICATION NUMBER: 60/362,158
PRIOR FILING DATE: 2002-03-05
NUMBER OF SEQ ID NOS: 455
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 42
LENGTH: 298
TYPE: PRT
ORGANISM: Homo sapiens
US-10-205-823-42

Query Match 32.1%; Score 486; DB 15; Length 298;
Best Local Similarity 36.7%; Pred. No. 1.1e-41;
Matches 101; Conservative 52; Mismatches 112; Indels 10; Gaps 4;
QY 6 HSLHYLFMGASODLGLSLFEALGYDDQLFVYDHSRRVPRTPWSSRISQWLQL 65
DB 28 YSLTYITGLSKHVEDVPAPFQALGSLNDLQFFRYNSKDRKSQPMGLWRQVE-GMEDWKQD 86
QY 66 SQLKGDHMTVDFTWIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDGDAL 125
DB 87 SQLQKAREDIFMETLKDIVEYNDNGSHVLOQRFCEIENNRSSGAFWKYYDGDYIE 146
QY 126 FCPDITLDMRAAPRAWPMTKLEWRHKIRARONRAYLERDCPAQLQQLLELGRGVLDQV 185
DB 147 FNKEIPAWVPDPAQITKQWEAEVYVQRAKAYLEECFATLRKYLKYSKNILDRQDP 206
QY 186 PLVKVT-HHVTSSVTLRCALNYYPQNTIMKWLKDKQPMDAKBEFPK----DVLPGDGT 241
DB 207 PSVVVTSQAPGKXKLCCLAYDFYPKIDVHWTAGEVQ-----BPRLRGDLVHNGNGT 261
QY 242 YQGWITLAVPGEQRYTCQVEHPGLDPLIWIWE 276
DB 262 YQSWVVAVPPQDTAPYSCHVQHSSLAQPLVVPWE 296

RESULT 10
US-10-177-293-23
Sequence 23, Application US/10177293
Publication No. US20030124128A1
GENERAL INFORMATION:
APPLICANT: Lilie, James
APPLICANT: Glatt, Karen
APPLICANT: Zhao, Xumei
APPLICANT: Gannavarpu, Manjula
APPLICANT: Kamatkar, Shubhangi
APPLICANT: Mertens, Maureen
APPLICANT: Myer, Vic
APPLICANT: Wang, Youzhen
APPLICANT: Xu, Yongyao
APPLICANT: Hoersch, Sebastian
APPLICANT: Monahan, John
APPLICANT: Meyers, Rachel E.
APPLICANT: Bast Jr., Robert C.
APPLICANT: Hortobagyi, Gabriel N.
APPLICANT: Pusztai, Lajos
APPLICANT: Meric, Funda
APPLICANT: Sahin, Aysegul

APPLICANT: Mills, Gordon B.
TITLE OF INVENTION: COMPOSITIONS, KITS, AND METHODS FOR IDENTIFICATION, ASSESSMENT,
PREVENTION, AND THERAPY OF BREAST CANCER
FILE REFERENCE: MRI-038
CURRENT APPLICATION NUMBER: US/10/177,293
CURRENT FILING DATE: 2002-06-21
PRIOR APPLICATION NUMBER: US 60/299,887
PRIOR FILING DATE: 2001-06-21
PRIOR APPLICATION NUMBER: US 60/301,572
PRIOR FILING DATE: 2001-06-27
PRIOR APPLICATION NUMBER: US 60/306,501
PRIOR FILING DATE: 2001-07-18
PRIOR APPLICATION NUMBER: US 60/325,002
PRIOR FILING DATE: 2001-09-25
PRIOR APPLICATION NUMBER: US 60/362,585
PRIOR FILING DATE: 2002-03-05
PRIOR APPLICATION NUMBER: US 60/xxx,xxx
PRIOR FILING DATE: 2002-05-14
NUMBER OF SEQ ID NOS: 506
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 23
LENGTH: 298
TYPE: PRT
ORGANISM: Homo sapiens
US-10-177-293-23

Query Match 32.1%; Score 486; DB 15; Length 298;
Best Local Similarity 36.7%; Pred. No. 1.1e-41;
Matches 101; Conservative 52; Mismatches 112; Indels 10; Gaps 4;
QY 6 HSLHYLFMGASODLGLSLFEALGYDDQLFVYDHSRRVPRTPWSSRISQWLQL 65
DB 28 YSLTYITGLSKHVEDVPAPFQALGSLNDLQFFRYNSKDRKSQPMGLWRQVE-GMEDWKQD 86
QY 66 SQLKGDHMTVDFTWIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDGDAL 125
DB 87 SQLQKAREDIFMETLKDIVEYNDNGSHVLOQRFCEIENNRSSGAFWKYYDGDYIE 146
QY 126 FCPDITLDMRAAPRAWPMTKLEWRHKIRARONRAYLERDCPAQLQQLLELGRGVLDQV 185
DB 147 FNKEIPAWVPDPAQITKQWEAEVYVQRAKAYLEECFATLRKYLKYSKNILDRQDP 206
QY 186 PLVKVT-HHVTSSVTLRCALNYYPQNTIMKWLKDKQPMDAKBEFPK----DVLPGDGT 241
DB 207 PSVVVTSQAPGKXKLCCLAYDFYPKIDVHWTAGEVQ-----BPRLRGDLVHNGNGT 261
QY 242 YQGWITLAVPGEQRYTCQVEHPGLDPLIWIWE 276
DB 262 YQSWVVAVPPQDTAPYSCHVQHSSLAQPLVVPWE 296

RESULT 11
US-10-015-535-32
Sequence 32, Application US/10015535
Publication No. US20030036506A1
GENERAL INFORMATION:
APPLICANT: Kranz, David M.
APPLICANT: Brophy, Susan
TITLE OF INVENTION: Mutated Class I Major Histocompatibility proteins and
Complexes
FILE REFERENCE: 100-00
CURRENT APPLICATION NUMBER: US/10/015,535
CURRENT FILING DATE: 2001-12-10
PRIOR APPLICATION NUMBER: 60/254,495
PRIOR FILING DATE: 2000-12-08
NUMBER OF SEQ ID NOS: 37
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 32
LENGTH: 542
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Synthetic


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Qy 182 QVPPPLVKKVTHV-TSSVTTLCRCALNYYPNITMKWLKQKQPMDAKEFPKQVLPNGDG 240
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 234 RTDSPKAVHTHSRSPEDKVTLCRCWALGFYPADITLTWQNGEEL-IQDMELVETRPAJDG 298
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Qy 241 TTQGWITLAVPGBEQRYTCQVEHPGLDQPLIWIWE 276
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
Db 293 TFQKASVVVPLGKEQYVTCVYHQGLPEPLTLRWE 328
      ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

Search completed: August 5, 2003, 13:21:56
Job time : 34 secs

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; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; ; OTHER INFORMATION: peptide
US-10-015-535-22

Query Match          30.9%; Score 468; DB 15; Length 540;
Best Local Similarity 39.5%; Pred. No. 1.7e-39;
Matches 109; Conservative 39; Mismatches 120; Indels 8; Gaps 7/

QY      6 HSLHLEPMGASEQDLGLSLFEALGYDDQLPVFYDH--ESRRVPEPTPWVSSRISQMWL 63
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db      144 HSLRYFYTAVSRPGLGEPRYMEVGYVDTEFVRFDSDAENRVEYEPRAWRMEQE-GPEYWE 202

QY      64 QLSQSLKGDHMFVTDFWTIMENHNASK-ESHTLQVILGCEWQEDNS--TEGYWKYGYDQG 121
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db      203 RETQKAKGNQSQSFVRDLRTLLGYYNQSGQSHTTQVTSIGCEVSGDGRLLRGYQQYAYDGC 262

QY      122 DALEFCDPTLDWAAEPRAMPFTKLEWRHKTRARQNRAYLERDCAQLOQLLELGRGVL D 181
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db      263 DYTALNEDLKTWTAADMAALITKHKWEQAG-EAEELRAYLEGTQVEWLRRYLKKGNA TLL 321

QY      182 QQVPLPVKVTHV--TSSVTTLRCALNYYPQNTIMKWLKKQPMDAKEFEFKDVLNPDG D 240
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db      322 RTDSPRAHVTHHSRPEDKVLTLCWALGFYPADIITLTQLNGEEL-IQDMELVETRPAGD G 380

QY      241 TYQGWITLAVPPGEEQRYTCQVHPGLDQPLIVIWE 276
      ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
Db      381 TFOKASVVPVLGKEQYTTCHVHQGLPELPITLWE 416
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RESULT 15
US-10-015-535-28
; Sequence 28, Application US/10015535
; Publication No. US20030036506A1
; GENERAL INFORMATION:
; APPLICANT: Kranz, David M.
; APPLICANT: Brophy, Susan
; TITLE OF INVENTION: Mutated Class I Major Histocompatibility proteins and
; FILE REFERENCE: 100-00
; CURRENT APPLICATION NUMBER: US/10/015,535
; CURRENT FILING DATE: 2001-12-10
; PRIOR APPLICATION NUMBER: 60/254,495
; PRIOR FILING DATE: 2000-12-08
; NUMBER OF SEQ ID NOS: 37
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 28
; LENGTH: 541
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; OTHER INFORMATION: peptide
US-10-015-535-28

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Query Match	30.9%	Score 468	DB 15	Length 541
Best Local Similarity	39.5%	Pred. No. 1.7e-39		
Matches	109	Conservative 39	Mismatches 120	Indels 8
Gaps	7			
QY	6	HSLSHLYFMGASEODLGLSLFEALGYDDQLFVFYDH--ESRRVPEPRPTVSSRISOQWVL	63	
Db	56	HSLSRYFTVAISRPLGSEPRYMEGVYDDTFVFRFDSDAENPRYEPFRARWMEQE-GPEYWE	114	
QY	64	QLSLSLKGMDHMTVDVFTWMENHNASK-ESHTLQVILGCEMQEDNS-TEGYWKYGYDQG	121	
Db	115	RETQKAKGNEQSFRLDLRTLILGYNYSKSGSHTIQVLSGCEVSDGRLLRGYQYAYDGC	174	
QY	122	DALEFCPTDLTDRAASPRAWPTKLEWRHKIRARONRAYLERDCAQLQQLLELGRGVLD	181	
Db	175	DYIALMEDIKTWTAAADMAALITKHKEWQAG-EAERLRAYLEGTCEVWLRYRYKNGNATLL	233	

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:07:04 ; Search time 14.5 Seconds
(without alignments)
805.365 Million cell updates/sec

Title: US-10-092-404-3
Perfect score: 1514
Sequence: 1 RLRSLSHLFLWGASEQDL.....RYTCQVHPLDQPLIVWE 276

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 4231058 residues

Total number of hits satisfying chosen parameters: 328717

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents AA:*
1: /cgn2_6/prodata/1/iaa/5A COMB.pep.*
2: /cgn2_6/prodata/1/iaa/5B COMB.pep.*
3: /cgn2_6/prodata/1/iaa/6A COMB.pep.*
4: /cgn2_6/prodata/1/iaa/6B COMB.pep.*
5: /cgn2_6/prodata/1/iaa/PCTUS COMB.pep.*
6: /cgn2_6/prodata/1/iaa/backfiles1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1514	100.0	276	4	US-09-094-964-3
2	1502	99.2	276	4	US-09-094-964-1
3	1502	99.2	348	3	US-08-652-265-2
4	1502	99.2	348	3	US-08-834-497A-2
5	1502	99.2	348	3	US-09-503-444A-2
6	1502	99.2	348	4	US-09-277-457-2
7	1502	99.2	348	4	US-09-679-729-2
8	1493	98.6	276	4	US-09-094-964-2
9	1493	98.6	348	3	US-08-652-265-6
10	1493	98.6	348	3	US-08-834-497A-6
11	1491	98.6	348	3	US-09-503-444A-6
12	1491	98.5	348	3	US-08-652-265-4
13	1491	98.5	348	3	US-08-834-497A-4
14	1491	98.5	348	3	US-09-503-444A-4
15	1482	97.9	348	3	US-08-652-265-8
16	1482	97.9	348	3	US-08-834-497A-8
17	1482	97.9	348	3	US-09-503-444A-8
18	517	34.1	361	3	US-08-652-265-22
19	517	34.1	361	3	US-08-834-497A-22
20	517	34.1	361	3	US-09-503-444A-22
21	511	33.8	364	4	US-08-914-372C-11
22	508	33.6	365	3	US-08-652-265-23
23	508	33.6	365	3	US-08-834-497A-23
24	508	33.6	365	3	US-09-503-444A-23
25	500	33.0	274	2	US-08-484-905-107
26	500	33.0	274	3	US-08-481-985B-107
27	500	33.0	274	3	US-08-370-476-107

28	500	33.0	341	3	US-08-890-719-38	Sequence 38, Appl
29	499	33.0	365	2	US-08-484-905-97	Sequence 97, Appl
30	499	33.0	365	3	US-08-481-985B-97	Sequence 97, Appl
31	499	33.0	365	3	US-08-370-476-97	Sequence 97, Appl
32	498	32.9	274	2	US-08-484-905-108	Sequence 108, App
33	498	32.9	274	3	US-08-481-985B-108	Sequence 108, App
34	498	32.9	274	3	US-08-370-476-108	Sequence 108, App
35	498	32.9	365	2	US-08-484-905-100	Sequence 100, App
36	498	32.9	365	3	US-08-481-985B-100	Sequence 100, App
37	498	32.9	365	3	US-08-370-476-100	Sequence 100, App
38	497	32.8	274	1	US-08-222-851-1	Sequence 1, Appli
39	497	32.8	363	4	US-08-914-372C-37	Sequence 37, Appl
40	497	32.8	365	2	US-08-484-905-99	Sequence 99, Appl
41	497	32.8	365	3	US-08-481-985B-99	Sequence 99, Appl
42	497	32.8	365	3	US-08-370-476-99	Sequence 99, Appl
43	496	32.8	274	2	US-08-484-905-106	Sequence 106, App
44	496	32.8	274	3	US-08-481-985B-106	Sequence 106, App
45	496	32.8	274	3	US-08-370-476-106	Sequence 106, App

ALIGNMENTS

RESULT 1
US-09-094-964-3
; Sequence 3, Application US/09094964
; Patent No. 6391852
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; APPLICANT: Bjorkman, Pamela J.
; APPLICANT: Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; TITLE OF INVENTION: AND IRON DEFICIENCY DISEASES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 276 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; US-09-094-964-3

Query Match 100.0%; Score 1514; DB 4; Length 276;
Best Local Similarity 100.0%; Pred. No. 6.2e-144;

	Matches	276;	Conservative	0;	Mismatches	0;	Indels	0;	Gaps	0;
Qy	1	RLLRSHSLHYLFMGASBODLGLSLFEALGYVDDQLFVFYDHESRRVEPRTPWSSRISSQ	60							
Db	1	RLLRSHSLHYLFMGASBODLGLSLFEALGYVDDQLFVFYDHESRRVEPRTPWSSRISSQ	60							
Qy	61	MLQLSLSKGDHMFVTVDFTMINENNASKESTLQVLGCMEONSTEGYWKYGVDG	120							
Db	61	MLQLSLSKGDHMFVTVDFTMINENNASKESTLQVLGCMEONSTEGYWKYGVDG	120							
Qy	121	QDALEFCPDTLDWRAAEPRAWPTTKLEWERHKIRARQNRAYLERDCAQLQQLLBLGRGVL	180							
Db	121	QDALEFCPDTLDWRAAEPRAWPTTKLEWERHKIRARQNRAYLERDCAQLQQLLBLGRGVL	180							
Qy	181	DQGVPLPVKYTHHTVSSTTLRCCALANYYPONITMKWLKKQPMDAKEFEFKDVLPLNGDG	240							
Db	181	DQGVPLPVKYTHHTVSSTTLRCCALANYYPONITMKWLKKQPMDAKEFEFKDVLPLNGDG	240							
Qy	241	TYCGWITLAVPPGEEORYTCOVERHPGLDQPLVIWE	276							
Db	241	TYCGWITLAVPPGEEORYTCOVERHPGLDQPLVIWE	276							

RESULT 2
 US-09-094-964-1
 ; Sequence 1, Application US/09094964
 ; Patent No. 6391852
 ; GENERAL INFORMATION:
 ; APPLICANT: Feder, John N.
 ; APPLICANT: Bjorkman, Pamela J.
 ; APPLICANT: Schatzman, Randall C.
 ; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
 ; TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
 ; TITLE OF INVENTION: AND IRON DEFICIENCY DISEASES
 ; NUMBER OF SEQUENCES: 5
 ; CORRESPONDENCE ADDRESS:
 ; ADDRESSEE: Pennie & Edmonds, LLP
 ; STREET: 1155 Avenue of the Americas
 ; CITY: New York
 ; STATE: NY
 ; COUNTRY: USA
 ; ZIP: 10036-2811
 ; COMPUTER READABLE FORM:
 ; MEDIUM TYPE: Diskette
 ; COMPUTER: IBM Compatible
 ; OPERATING SYSTEM: Windows
 ; SOFTWARE: FastSeq for Windows Version 2.0b
 ; CURRENT APPLICATION DATA:
 ; APPLICATION NUMBER: US/09/094,964
 ; FILING DATE: June 12, 1998
 ; CLASSIFICATION:
 ; PRIOR APPLICATION DATA:
 ; APPLICATION NUMBER: 08/876,010
 ; FILING DATE: June 13, 1997
 ; ATTORNEY/AGENT INFORMATION:
 ; NAME: Poissant, Brian M
 ; REGISTRATION NUMBER: 28,462
 ; REFERENCE/DOCKET NUMBER: 8907-0074-999
 ; TELECOMMUNICATION INFORMATION:
 ; TELEPHONE: 650-493-4935
 ; TELEFAX: 650-493-5556
 ; TELEX: 66141 PENNIE
 ; INFORMATION FOR SEQ ID NO: 1:
 ; SEQUENCE CHARACTERISTICS:
 ; LENGTH: 276 amino acids
 ; TYPE: amino acid
 ; STRANDEDNESS: single
 ; TOPOLOGY: linear
 ; MOLECULE TYPE: peptide
 US-09-094-964-1

Matches 274; Conservative 0; Mismatches 2; Indels	
Qy	1 RLLRSHSLHYLFMGASBQDGLGLSLFEALGYVDDOLFVYDHESRRVPRPT
Db	1 RLLRSHSLHYLFMGASBQDGLGLSLFEALGYVDDOLFVYDHESRRVPRPT
Qy	61 MWLQLSQSLKGWDHMFVTVDFTWIMENHNASKESHTLVILGCENQDNST
Db	61 MWLQLSQSLKGWDHMFVTVDFTWIMENHNASKESHTLVILGCENQDNST
Qy	121 QDALEFCPDTLDWRAAPRAWPTKLEWERHKIRARQNRAVLERDCCPAQLQ
Db	121 QDALEFCPDTLDWRAAPRAWPTKLEWERHKIRARQNRAVLERDCCPAQLQ
Qy	181 DQQVPLPVKTVHHVTSSTVTTLRCRAlNYPQNIITMKWLKDKQPMDAKEFE
Db	181 DQQVPLPVKTVHHVTSSTVTTLRCRAlNYPQNIITMKWLKDKQPMDAKEFE
Qy	241 TYQGWITLAVPPGSEORYTCQVEHPGLDPLIVWE 276
Db	241 TYQGWITLAVPPGSEORYTCQVEHPGLDPLIVWE 276
RESULT 3	
US-08-652-265-2	
; Sequence 2, Application US/08652265	
; Patent No. 6025130	
; GENERAL INFORMATION:	
; APPLICANT: Thomas, Winston J.	
; APPLICANT: Drayna, Dennis T.	
; APPLICANT: Feder, John N.	
; APPLICANT: Gnirke, Andreas	
; APPLICANT: Ruddy, David	
; APPLICANT: Tsuchihashi, Zenta	
; APPLICANT: Wolff, Roger K.	
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene	
; NUMBER OF SEQUENCES: 44	
; CORRESPONDENCE ADDRESSES:	
; ADDRESSEE: Townsend and Townsend and Crew LLP	
; STREET: Two Embarcadero Center, Eighth Floor	
; CITY: San Francisco	
; STATE: California	
; COUNTRY: USA	
; ZIP: 94111-3834	
; COMPUTER READABLE FORM:	
; MEDIUM TYPE: Floppy disk	
; COMPUTER: IBM PC compatible	
; OPERATING SYSTEM: PC-DOS/MS-DOS	
; SOFTWARE: PatentIn Release #1.0, Version #1.30	
; CURRENT APPLICATION DATA:	
; APPLICATION NUMBER: US/08/652,265	
; FILING DATE: 23-MAY-1996	
; CLASSIFICATION: 514	
; ATTORNEY/AGENT INFORMATION:	
; NAME: Smith, William M.	
; REGISTRATION NUMBER: 30,223	
; REFERENCE/DOCKET NUMBER: 17957-000500	
; TELECOMMUNICATION INFORMATION:	
; TELEPHONE: (415) 576-0200	
; TELEFAX: (415) 576-0300	
; INFORMATION FOR SEQ ID NO: 2:	
; SEQUENCE CHARACTERISTICS:	
; LENGTH: 348 amino acids	
; TYPE: amino acid	
; TOPOLOGY: linear	
; MOLECULE TYPE: protein	
US-08-652-265-2	

Query Match 99.2%; Score 1502; DB 4; Length 276;
 Best Local Similarity 99.3%; Pred. No. 9.9e-143;

Db 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDQDLFFVYDHSRRRVEPRTPWVSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
Db 83 MWLQLSQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALFECPTDLDWRAAEPRAMPKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
Db 143 QDHLFECPTDLDWRAAEPRAMPKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DOQVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQMDAKEPEPKDVLPGDG 240
Db 203 DOQVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQMDAKEPEPKDVLPGDG 262
QY 241 TYQGWITLAVPPGEQRQYTCQVEHPGLDQPLIVWE 276
Db 263 TYQGWITLAVPPGEQRQYTCQVEHPGLDQPLIVWE 298

RESULT 4

US-08-834-497A-2
; Sequence 2, Application US/08834497A
; Patent No. 6140305

; GENERAL INFORMATION:

; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
; NUMBER OF SEQUENCES: 76
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036-2811

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: FASTSEQ for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/834,497A
; FILING DATE: 04-APR-1997

; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 08/632,673
; FILING DATE: 16-APR-1996
; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 08/630,912
; FILING DATE: 04-APR-1996
; CLASSIFICATION: 514

; ATTORNEY/AGENT INFORMATION:

; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0056-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids

; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-834-497A-2

Query Match

Best Local Similarity 99.2%; Score 1502; DB 3; Length 348;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDQDLFFVYDHSRRRVEPRTPWVSSRISSQ 60
Db 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDQDLFFVYDHSRRRVEPRTPWVSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
Db 83 MWLQLSQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALFECPTDLDWRAAEPRAMPKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
Db 143 QDHLFECPTDLDWRAAEPRAMPKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DOQVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQMDAKEPEPKDVLPGDG 240
Db 203 DOQVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQMDAKEPEPKDVLPGDG 262
QY 241 TYQGWITLAVPPGEQRQYTCQVEHPGLDQPLIVWE 276
Db 263 TYQGWITLAVPPGEQRQYTCQVEHPGLDQPLIVWE 298

RESULT 5

US-09-503-444A-2

; Sequence 2, Application US/09503444A
; Patent No. 6228594

; GENERAL INFORMATION:

; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: WordPerfect Version 8
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/503,444A
; FILING DATE: 14-Feb-2000

; CLASSIFICATION:

; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/652,265
; FILING DATE: 23-May-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,673
; FILING DATE: 16-Apr-1996

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: 08/630,912
; FILING DATE: 04-Apr-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0088-999

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-790-9090

TELEFAX: 212-869-9741

TELEX: 66141

INFORMATION FOR SEQ ID NO: 2:

SEQUENCE CHARACTERISTICS:

LENGTH: 348 amino acids

TYPE: amino acid

TOPOLOGY: linear

MOLECULE TYPE: protein

US-09-503-444A-2

Query Match 99.2%; Score 1502; DB 3; Length 348;

Best Local Similarity 99.3%; Pred. No. 1.4e-142;

Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPTPWSSRISSQ 60

Db 23 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPTPWSSRISSQ 82

QY 61 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120

Db 83 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142

QY 121 QDALEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDPCPAQLQQLLELGRGVL 180

Db 143 QDHLEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDPCPAQLQQLLELGRGVL 202

QY 181 DQVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240

Db 203 DQVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 262

QY 241 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 276

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 181 DQVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240

Db 203 DQVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 262

QY 241 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 276

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

Db 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

QY 263 TYQGWITLAVPPEEQRVTCQVEHGLDQPLIVWE 298

RESULT 8

US-09-094-964-2

; Sequence 2, Application US/09094964

; Patent No. 6391852

; GENERAL INFORMATION:

; APPLICANT: Feder, John N.

; APPLICANT: Bjorkman, Pamela J.

; APPLICANT: Schatzman, Randall C.

; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR

; TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES

; NUMBER OF SEQUENCES: 5

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Pennie & Edmonds, LLP

; STREET: 1155 Avenue of the Americas

; CITY: New York

; STATE: NY

RESULT 8

US-09-094-964-2

; Sequence 2, Application US/09094964

; Patent No. 6391852

; GENERAL INFORMATION:

; APPLICANT: Feder, John N.

; APPLICANT: Bjorkman, Pamela J.

; APPLICANT: Schatzman, Randall C.

; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR

; TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES

; NUMBER OF SEQUENCES: 5

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Pennie & Edmonds, LLP

; STREET: 1155 Avenue of the Americas

; CITY: New York

; STATE: NY

COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows
SOFTWARE: FASTSEQ for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/094,964
FILING DATE: June 12, 1998
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/876,010
FILING DATE: June 13, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0074-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-094-964-2

Query Match 98.6%; Score 1493; DB 4; Length 276;

Best Local Similarity 98.9%; Pred. No. 7.9e-142;

Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 60
QY 61 MWLQSLQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQSLQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDALEFCPTDLDWRAAPRAWPTKLEWERHKIRARQRAYLERDPCPAQLQQLLELGRGVL 180
DB 121 QDHLFCPTDLDWRAAPRAWPTKLEWERHKIRARQRAYLERDPCPAQLQQLLELGRGVL 180
QY 181 DQOVPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 181 DQOVPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
QY 241 TYQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 276
DB 241 TYQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 276

RESULT 9

US-08-652-265-6

Sequence 6, Application US/08652265

Patent No. 6025130

GENERAL INFORMATION:

APPLICANT: Thomas, Winston J.

APPLICANT: Drayna, Dennis T.

APPLICANT: Feder, John N.

APPLICANT: Gnirke, Andreas

APPLICANT: Ruddy, David

APPLICANT: Tsuchihashi, Zenta

APPLICANT: Wolff, Roger K.

TITLE OF INVENTION: Hereditary Hemochromatosis Gene

NUMBER OF SEQUENCES: 44

CORRESPONDENCE ADDRESS:

ADDRESSEE: Townsend and Townsend and Crew LLP

STREET: Two Embarcadero Center, Eighth Floor

CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94111-3834
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Smith, William M.
REGISTRATION NUMBER: 30,223
REFERENCE/DOCKET NUMBER: 17957-000500
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 576-0200
TELEFAX: (415) 576-0300
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-652-265-6

Query Match 98.6%; Score 1493; DB 3; Length 348;

Best Local Similarity 98.9%; Pred. No. 1.1e-141;

Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 82
QY 61 MWLQSLQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPTDLDWRAAPRAWPTKLEWERHKIRARQRAYLERDPCPAQLQQLLELGRGVL 180
DB 143 QDHLFCPTDLDWRAAPRAWPTKLEWERHKIRARQRAYLERDPCPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 203 DQOVPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 262
QY 241 TYQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 298

RESULT 10

US-08-834-497A-6

Sequence 6, Application US/08834497A

Patent No. 6140305

GENERAL INFORMATION:

APPLICANT: Thomas, Winston J.

APPLICANT: Drayna, Dennis T.

APPLICANT: Feder, John N.

APPLICANT: Gnirke, Andreas

APPLICANT: Ruddy, David

APPLICANT: Tsuchihashi, Zenta

APPLICANT: Wolff, Roger K.

TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS

NUMBER OF SEQUENCES: 76

CORRESPONDENCE ADDRESS:

ADDRESSEE: pennie & Edmonds LLP

STREET: 1155 Avenue of the Americas

CITY: New York

STATE: New York

COUNTRY: USA

ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FASTSEQ for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/834,497A
FILING DATE: 04-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/632,673
FILING DATE: 16-APR-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/630,912
FILING DATE: 04-APR-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0056-999
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-834-497A-6

Query Match 98.6%; Score 1493; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.1e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYDDQLFVFDHESRRVPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASEQDLGLSLFEALGYDDQLFVFDHESRRVPRTPWVSSRISSQ 82
QY 61 MWLQSLQSLKGWDMFTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDMFTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 ODALFPCPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGNGD 240
DB 203 DQVPPPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGNGD 262
QY 241 TYQGWITLAVPGEQRYTCQVEHPGLDQPLIWIWE 276
DB 263 TYQGWITLAVPGEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 11

US-09-503-444A-6
Sequence 6, Application US/09503444A
Patent No. 6228594
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David

APPLICANT: Teuchiashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Hereditary Hemochromatosis Gene
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: WordPerfect Version 8
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/503,444A
FILING DATE: 14-Feb-2000
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/652,265
FILING DATE: 23-May-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/632,673
FILING DATE: 16-Apr-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/630,912
FILING DATE: 04-Apr-1996
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0088-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-790-9090
TELEFAX: 212-869-9741
TELEX: 66141
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-503-444A-6
Query Match 98.6%; Score 1493; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.1e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYDDQLFVFDHESRRVPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASEQDLGLSLFEALGYDDQLFVFDHESRRVPRTPWVSSRISSQ 82
QY 61 MWLQSLQSLKGWDMFTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDMFTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 ODALFPCPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGNGD 240
DB 203 DQVPPPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGNGD 262
QY 241 TYQGWITLAVPGEQRYTCQVEHPGLDQPLIWIWE 276
DB 263 TYQGWITLAVPGEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 12

US-08-652-265-4
Sequence 4, Application US/08652265

Patent No. 6025130
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Hereditary Hemochromatosis Gene
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend and Crew LLP
STREET: Two Embarcadero Center, Eighth Floor
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94111-3834
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Smith, William M.
REGISTRATION NUMBER: 30,223
REFERENCE/DOCKET NUMBER: 17957-000500
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 576-0200
TELEFAX: (415) 576-0300
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-652-265-4

Query Match 98.5%; Score 1491; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.7e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISQ 60
DB 23 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISQ 82
QY 61 MWLQSLQSLKGWDHMTVDFTWMENHNHNSKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDHMTVDFTWMENHNHNSKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGYL 180
DB 143 QDHLFCPDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGYL 202
QY 181 DOQVPLVKVTHHTVSSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
DB 203 DOQVPLVKVTHHTVSSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 262
QY 241 TYQGWITLAVPGEQRQYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPGEQRQYTCQVEHPGLDQPLIVWE 298

RESULT 13

US-08-834-497A-4
Sequence 4, Application US/08834497A
Patent No. 6140305
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.

APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FastSeq for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/834,497A
FILING DATE: 04-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
APPLICATION NUMBER: US 08/632,673
FILING DATE: 16-APR-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/630,912
FILING DATE: 04-APR-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0056-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-834-497A-4
Query Match 98.5%; Score 1491; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.7e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISQ 60
DB 23 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISQ 82
QY 61 MWLQSLQSLKGWDHMTVDFTWMENHNHNSKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDHMTVDFTWMENHNHNSKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGYL 180
DB 143 QDHLFCPDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGYL 202
QY 181 DOQVPLVKVTHHTVSSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
DB 203 DOQVPLVKVTHHTVSSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 262
QY 241 TYQGWITLAVPGEQRQYTCQVEHPGLDQPLIVWE 276

Db 263 TYQGWITLAVPGEQRYTQVEHPGLDQPLIVWE 298
|||||
RESULT 14
US-09-503-444A-4
; Sequence 4, Application US/09503444A
; Patent No. 6228594
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: WordPerfect Version 8
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/503,444A
; FILING DATE: 14-Feb-2000
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/652,265
; FILING DATE: 23-May-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,673
; FILING DATE: 16-Apr-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/630,912
; FILING DATE: 04-Apr-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0088-999
; TELEPHONE: 212-790-9090
; TELEFAX: 212-869-9741
; TELEX: 66141
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-09-503-444A-4
Query Match 98.5%; Score 1491; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.7e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFPMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISSQ 60
Db 23 RLLRSHSLHYLFPMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISSQ 82
QY 61 MWLQLSQSLKGWDHMTFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
Db 83 MWLQLSQSLKGWDHMTFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGYL 180
Db 143 QHLEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGYL 202
QY 181 DOQVPLVKVTHVTSSVTTLCRALNYYPQNTWKWLKDKQPMDAKEPEPKDVLPGDVG 240

Db 143 ODHLEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGYL 202
QY 181 DOQVPLVKVTHVTSSVTTLCRALNYYPQNTWKWLKDKQPMDAKEPEPKDVLPGDVG 240
Db 203 DOQVPLVKVTHVTSSVTTLCRALNYYPQNTWKWLKDKQPMDAKEPEPKDVLPGDVG 262
QY 241 TYQGWITLAVPGEQRYTQVEHPGLDQPLIVWE 276
Db 263 TYQGWITLAVPGEQRYTQVEHPGLDQPLIVWE 298
RESULT 15
US-08-652-265-8
; Sequence 8, Application US/08652265
; Patent No. 6025130
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M.
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 17957-000500
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-652-265-8
Query Match 97.9%; Score 1482; DB 3; Length 348;
Best Local Similarity 98.6%; Pred. No. 1.4e-140;
Matches 272; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
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Db 23 RLLRSHSLHYLFPMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISSQ 82
QY 61 MWLQLSQSLKGWDHMTFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
Db 83 MWLQLSQSLKGWDHMTFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGYL 180
Db 143 QHLEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGYL 202
QY 181 DOQVPLVKVTHVTSSVTTLCRALNYYPQNTWKWLKDKQPMDAKEPEPKDVLPGDVG 240

Search completed: August 5, 2003, 13:11:12
Job time : 15.5 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:05:29 ; Search time 38 Seconds
(without alignments)

1152.856 Million cell updates/sec

Title: US-10-092-404-2

Perfect score: 1520

Sequence: 1 RLRLSHLHLYFMGASEQDL.....RYTCQVHPGLDPLIVWE 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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24: /SIDS1/gcgdata/geneseq/geneseq-emb1/AA2003.DAT.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1520	100.0	276	20 AAW94296	HFE mutant (H63D-H
2	1520	100.0	276	24 ABG72686	Human haemochromat
3	1520	100.0	348	22 AAB36871	Human hereditary h
4	1513	99.5	276	20 AAW94295	Wild-type HFE poly
5	1513	99.5	276	24 ABG72685	Human haemochromat
6	1513	99.5	348	18 AAW36499	Hereditary haemoch
7	1513	99.5	348	21 AAB19149	A human histocoma
8	1513	99.5	348	22 AAB36869	Human hereditary h
9	1509	99.3	348	22 AAB36872	Human hereditary h

10	1508	99.2	438	23 AAW00035	Beta 2 microglobul
11	1502	98.8	348	22 AAB36870	Human hereditary h
12	1493	98.2	276	20 AAW94297	HFE mutant (H111A/
13	1493	98.2	276	24 ABG72687	Human haemochromat
14	523	34.4	361	22 AAB36873	Rabbit leukocyte a.
15	514	33.8	92	24 ABF68379	Human colon specif
16	514	33.8	365	22 AAB36874	MHC class I protei
17	506	33.3	274	21 AAY68275	Human leukocyte an
18	506	33.3	274	21 AAY52929	HLA-A2/A28 family
19	506	33.3	274	22 AAB58690	HLA-A2/A28 protein
20	506	33.3	280	22 ABU10225	Human leukocyte an
21	506	33.3	280	24 ABU08672	Human histocompat
22	506	33.3	415	22 ABU10224	Human partial beta
23	506	33.3	415	24 ABU08671	Human single chain
24	505	33.2	365	21 AAY68265	Human leukocyte an
25	505	33.2	365	21 AAY52919	HLA-A2/A28 family
26	505	33.2	365	22 AAB58680	Human leukocyte an
27	505	33.2	368	22 AAM24017	Human EST encoded
28	504	33.2	274	21 AAY68276	Human leukocyte an
29	504	33.2	274	21 AAY52930	HLA-A2/A28 family
30	504	33.2	274	22 AAB58691	HLA-A2/A28 protein
31	504	33.2	365	21 AAY68268	Human leukocyte an
32	504	33.2	365	21 AAY52922	HLA-A2/A28 family
33	504	33.2	365	22 AAB58683	HLA-A2/A28 protein
34	503	33.1	274	9 AAP80911	Consensus sequence
35	503	33.1	365	21 AAY68267	Human leukocyte an
36	503	33.1	365	21 AAY52921	HLA-A2/A28 family
37	503	33.1	365	22 AAB58682	HLA-A2/A28 protein
38	502	33.0	274	21 AAY68274	Human leukocyte an
39	502	33.0	274	21 AAY52928	HLA-A2/A28 family
40	502	33.0	274	22 AAB58689	HLA-A2/A28 protein
41	502	33.0	365	21 AAY68266	Human leukocyte an
42	502	33.0	365	21 AAY52920	HLA-A2/A28 family
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44	501	33.0	412	19 AAW68385	Chimeric HLA-A2.1/
45	500	32.9	274	21 AAY68273	Human leukocyte an

ALIGNMENTS

RESULT 1

AAW94296

ID AAW94296 standard; peptide; 276 AA.

XX AC

XX AAW94296;

XX DT

27-APR-1999 (first entry)

XX DE

HFE mutant (H63D-HFE) polypeptide sequence.

XX DE

HFE; beta-2-microglobulin; beta2m; iron overload; hemochromatosis; transfusion; protein replacement therapy; hereditary hemochromatosis; transferrin receptor; iron deficiency; anemia; mutant.

XX OS

Synthetic.

XX FH

Key Location/Qualifiers

FT FT

Misc-difference 2 /note= "indicated in the sequence listing as Arg"

FT FT

Misc-difference 41

FT FT

/label= H63D

FT FT

/note= "wild type His (of the mature protein sequence) is replaced by Asp"

FT FT

W09856814-A1.

XX PN

17-DEC-1998.

XX PD

12-JUN-1998; 98WO-US12436.

XX PF

13-JUN-1997; 97US-0876010.

XX PR

AA036871
ID AAB36871 standard; Protein; 348 AA.

XX AC AAB36871;

XX DT 21-FEB-2001 (first entry)

XX DE Human hereditary hemochromatosis 24d2 mutation protein.

XX KW HH; hereditary hemochromatosis; chelation agent;

XX KW T-cell differentiation factor; iron overload.

XX OS Homo sapiens;

XX PN US6140305-A.

XX PD 31-OCT-2000.

XX PF 04-APR-1997; 97US-0834497.

XX PR 04-APR-1996; 96US-0630912.

XX PR 16-APR-1996; 96US-0632673.

XX PR 23-MAY-1996; 96US-0652265.

XX PA (BIRA) BIO-RAD LAB INC.

XX PI Thomas-WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;

XX PI Feder JN;

XX DR WPI; 2001-006341/01.

XX DR N-PSDB; AAC68427.

XX PT New hereditary hemochromatosis gene products or polypeptides, useful
PT for treating hereditary hemochromatosis in a patient, and as a metal
PT chelation agent alleviating iron overload -

XX PS Claim 3; Fig 4; 108pp; English.

XX CC The present invention relates to hereditary hemochromatosis gene
CC products. These proteins may be used to treat a patient diagnosed as
CC having human hemochromatosis disease. It is also useful as a metal
CC chelation agent or as a T-cell differentiation factor, and for
CC alleviating iron overload. They may also be used in protein replacement
CC therapy for individuals having a defective human hemochromatosis gene.

XX SQ Sequence 348 AA;

Query Match 100.0%; Score 1520; DB 22; Length 348;

Best Local Similarity 100.0%; Pred. No. 9.9e-135;

Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLIRSHSLHYLFMGASEQDGLSLFALGYDDQLFVYDDERRRVEPTPWSSRISQ 60

DB 23 RLIRSHSLHYLFMGASEQDGLSLFALGYDDQLFVYDDERRRVEPTPWSSRISQ 82

QY 61 MMLQLSLSKQWDHMTFVFTIMENHNHSHKESHTLQVILGCEMDESNSTEGYWKYGYDG 120

DB 83 MMLQLSLSKQWDHMTFVFTIMENHNHSHKESHTLQVILGCEMDESNSTEGYWKYGYDG 142

QY 121 QHLEFCPTDLWRAAPRAWPTKLEWRHKIRARONRAYLERDCAQQLLELGRGVL 180

DB 143 QHLEFCPTDLWRAAPRAWPTKLEWRHKIRARONRAYLERDCAQQLLELGRGVL 202

QY 181 DQOVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMADKEPEKDVLPNGDG 240

DB 203 DQOVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMADKEPEKDVLPNGDG 262

QY 241 TYOGWITLAVPPGEQRYTCQVEHPGLDQPLIIVIE 276

DB 263 TYOGWITLAVPPGEQRYTCQVEHPGLDQPLIIVIE 298

RESULT 4

AAW94295
ID AAW94295 standard; peptide; 276 AA.

XX AC AAW94295;

XX DT 27-APR-1999 (first entry)

XX DE Wild-type HFE polypeptide sequence.

XX KW HFE; beta-2-microglobulin; beta2m; iron overload; hemochromatosis;

XX KW transfusion; protein replacement therapy; hereditary hemochromatosis;

XX KW transferrin receptor; iron deficiency; anemia.

XX OS Unidentified.

XX FH Key Location/Qualifiers

XX FT Misc-difference 2

XX FT note= "indicated in the sequence listing as Arg"

XX PN W09856814-A1.

XX PD 17-DEC-1998.

XX PF 12-JUN-1998; 98WO-US12436.

XX PR 13-JUN-1997; 97US-0876010.

XX PA (CALY) CALIFORNIA INST OF TECHNOLOGY.

XX PA (PROG-) PROGENITOR INC.

XX PI Bjorkman PJ, Feder JN, Schatzman RC;

XX WPI; 1999-080886/07.

XX PT New treatment of an iron overload disease - comprises use of HFE

XX PT polypeptides provided in a complex with full length, wild type human

XX PT (2m), useful in protein replacement therapy

XX PS Claim 1; Page 13; 36pp; English.

XX CC The present sequence represents a wild-type HFE polypeptide. The HFE
XX polypeptides (AAW94295-297) provided in a complex with full length,
XX wild type human beta-2-microglobulin (beta2m) form compositions in the
XX treatment of primary iron overload diseases (e.g. hemochromatosis), or
XX other iron overload conditions resulting from secondary causes (e.g.
XX repeated transfusions). Data regarding the structure and function
XX correlations of HFE polypeptides is useful in designing drugs that
XX modulate the HFE gene and HFE activity. The polypeptides are also useful
XX in protein replacement therapy for individuals possessing a defective
XX HFE gene (e.g. Hereditary hemochromatosis). (Ant)agonists of the
XX polypeptides are also useful in treating primary and secondary iron
XX overload diseases. The modulators of the transferrin receptor are useful
XX in treating iron deficiency conditions such as anemia, and in modulating
XX the amount of iron transported into a cell. The HFE polypeptides provide
XX a molecular basis for the relationship between HFE and iron metabolism,
XX which enables treatment of iron overload and deficiency diseases.

XX SQ Sequence 276 AA;

Query Match 99.5%; Score 1513; DB 20; Length 276;

Best Local Similarity 99.6%; Pred. No. 3.3e-134;

Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLIRSHSLHYLFMGASEQDGLSLFALGYDDQLFVYDDERRRVEPTPWSSRISQ 60

DB 1 RLIRSHSLHYLFMGASEQDGLSLFALGYDDQLFVYDDERRRVEPTPWSSRISQ 60

QY 61 MMLQLSLSKQWDHMTFVFTIMENHNHSHKESHTLQVILGCEMDESNSTEGYWKYGYDG 120

DB 61 MMLQLSLSKQWDHMTFVFTIMENHNHSHKESHTLQVILGCEMDESNSTEGYWKYGYDG 120

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Db 121 QDHFPCPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
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Db 181 DQVPPPLVKVTHVTSSVTLRCRALNYYPONTMKWLKDKQPMDAKEPEPKDVLNPDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
Db 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 5
ABG72685
ID ABG72685 standard; protein; 276 AA.
XX
AC ABG72685;
XX
DT 05-MAR-2003 (first entry)
XX
DE Human haemochromatosis (HFE) mature protein.
XX
KW Human; haemochromatosis; HFE; hereditary haemochromatosis;
KW iron overload disease; iron deficiency disease; Beta2-microglobulin;
KW Beta2m; transferrin receptor; anaemia.
XX
OS Homo sapiens.
XX
PN US6391852-B1.
XX
PD 21-MAY-2002.
XX
PF 12-JUN-1998; 98US-0094964.
XX
PR 13-JUN-1997; 97US-0876010.
XX
PA (BIRA) BIO-RAD LAB INC.
PA (CALY) CALIFORNIA INST OF TECHNOLOGY.
XX
PI Feder JN, Bjorkman PJ, Schatzman RC;
XX
DR WPI; 2003-155377/15.
XX
XX
PT Method of treating an iron overload disease comprises administration of
PT a soluble complex comprising a 276 amino acid HFE polypeptide and a
PT full length, wild-type human beta2m -
XX
PS Claim 1; Column 1; 17pp; English.
XX
XX The invention relates to a method of treating an iron overload disease
CC comprising administration of a soluble complex comprising a 276 amino
CC acid mature HFE (hereditary haemochromatosis gene protein) polypeptide
CC (ABG72685-ABG72687) and a full length, wild-type human beta2m
CC (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of
CC "1,2-⁵¹I-transferrin in the presence of purified H63D-HFE/beta2m
CC heterodimers was determined. At a concentration of 250 nM H63D-HFE/
CC beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for
CC transferrin of 28 nM. At the same concentration of normal HFE/beta2m
CC heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence
CC of any HFE/beta2m heterodimers, TfR displayed a KD for transferrin of
CC 7nM. It was observed that H63D-HFE/beta2m heterodimers were 30-40 % less
CC efficient in decreasing TfR affinity for transferrin compared to
CC wild-type HFE. The method is useful for treating iron overload diseases
CC and iron deficiency e.g. anaemia. The present sequence is wild-type
CC mature HFE.
XX
SQ Sequence 276 AA;

Query Match 99.5%; Score 1513; DB 24; Length 276;
Best Local Similarity 99.6%; Pred. No. 3.3e-134;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISSQ 60
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Db 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISSQ 60
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Db 61 MWLQLSQSLSKGDHMTFTVDFWTIMENHNHSKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
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Db 181 DQVPPPLVKVTHVTSSVTLRCRALNYYPONTMKWLKDKQPMDAKEPEPKDVLNPDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
Db 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 6
AAW36499
ID AAW36499 standard; Protein; 348 AA.
XX
AC AAW36499;
XX
DT 14-APR-1998 (first entry)
XX
DE Hereditary haemochromatosis gene product.
XX
KW Hereditary haemochromatosis; metal toxicity; diagnosis;
KW gene therapy; prenatal screening; human.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT Misc-difference 63 /note= "substituted by Asp in 24s2 mutant"
FT Misc-difference 65 /note= "substituted by Cys in 24d7 variant"
FT Misc-difference 282 /note= "substituted by Tyr in 24d1 mutant"
FT
XX
PN WO9738137-A1.
XX
PD 16-OCT-1997.
XX
PF 04-APR-1997; 97WO-US06254.
XX
PR 23-MAY-1996; 96US-0652265.
PR 04-APR-1996; 96US-0630912.
PR 16-APR-1996; 96US-0632673.
XX
PA (MERC-) MERCATOR GENETICS INC.
XX
PI Drayna DT, Feder JN, Gnirke A, Ruddy D, Thomas WJ;
PI Tsuchihashi Z, Wolff RK;
XX
DR WPI; 1997-512743/47.
DR N-PSDB; AAT96690, AAT96691.
XX
PT Hereditary haemochromatosis gene and variants - useful for diagnosis
PT and treatment of hereditary haemochromatosis disease
XX
PS Disclosure; Fig 4; 115pp; English.
XX
CC This polypeptide is the expression product of a novel human gene
CC (see AAT96690) whose mutated form is associated with hereditary
CC haemochromatosis (HH). A single mutation (24d1) in the HH gene
CC appears responsible for the majority of HH disease. This comprises
CC a G to A substitution that is present in 86% of affected
CC chromosomes and in 4% of unaffected chromosomes. It results in a
CC Cys to Tyr substitution in the encoded protein at a critical
CC disulphide bridge important for secondary structure. The following

CC are claimed: the 10825 bp genomic DNA sequence (1), a 1437 bp cDNA
CC sequence (1a) (see AAR96691) and their 24d1, 24d2 and 24d7 variants;
CC a cloning or expression vector; host cells; a peptide product
CC chosen from the HH gene product, its variants (24d1, 24d2 and
CC 24d7), or a peptide of at least 56 amino acid residues of these; an
CC antibody produced using the peptide as an immunogen; a method to
CC determine the presence or absence of the common HH gene mutation;
CC an animal model for the HH disease; metal chelation agents, T-cell
CC differentiation factors and therapeutic agents for the mitigation
CC of injury due to oxidative processes in vivo or mitigation of iron
CC overload; a method for screening potential therapeutic agents for
CC activity in connection with HH disease; an antisense oligonucleotide
CC directed against a transcriptional product of a nucleic acid
CC sequence as above; and oligonucleotides or pairs of oligonucleotides
CC covering a range of nucleotides from (1), (1a) or their variants,
CC useful for detecting a polymorphism in the HH gene. The invention
CC also relates to methods for screening for HH homozygotes, to HH
CC diagnosis, prenatal screening and diagnosis, and therapies of HH
CC disease, including gene therapy, protein- and antibody-based
CC therapeutics, and small molecule therapeutics.

XX
SQ Sequence 348 AA;

Query Match 99.5%; Score 1513; DB 18; Length 348;
Best Local Similarity 99.6%; Pred. No. 4.5e-134;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISSQ 82
QY 61 MWLQSLQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCMQEDNSTEGYWKYGYDG 142
QY 121 QDHLEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNAYLERDCAQQLLELGRGVL 180
DB 143 QDHLEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNAYLERDCAQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSSVTLRCLALNYPQNTMKWLKDKQPMDAKEPEKDPVLPNGDG 240
DB 203 DQOVPLVKVTHVTSSVTLRCLALNYPQNTMKWLKDKQPMDAKEPEKDPVLPNGDG 262
QY 241 TYQGWTITLAVPPGEGORYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWTITLAVPPGEGORYTCQVEHPGLDQPLIVWE 298

RESULT 7
AAB19149
ID AAB19149 standard; Protein; 348 AA.
XX AAB19149;
XX
DT 19-FEB-2001 (first entry)
XX
DE A human histocompatibility iron loading (HFE) protein.
KW Human; histocompatibility iron loading protein; HFE protein;
KW major histocompatibility complex; non-classical class I gene;
KW chromosome 6p; iron disorder; haemochromatosis.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT Peptide 1..22
FT /note= "signal peptide"
FT Misc-difference 63
FT /note= "when nucleotide 187 is mutated to G, then
FT this residue is Asp"
FT Misc-difference 65
FT /note= "when nucleotide 193 is mutated to T, then
FT this residue is Cys"

FT Domain 90..108
FT Misc-difference 93 /note= "alpha1 domain"
FT /note= "when nucleotide 277 is mutated to C, then
FT this residue is Arg"
FT Misc-difference 105 /note= "when nucleotide 314 is mutated to C, then
FT this residue is Thr"
FT WO200058515-A1.
PN 05-OCT-2000.
XX
PD 24-MAR-2000; 2000WO-US07982.
XX
PF 26-MAR-1999; 99US-0277457.
XX
PR (BILL-) BILLUPS-ROTHENBERG INC.
XX
PI Rothenberg BE, Sawada-Hirai R, Barton JC;
XX
DR WPI; 2000-647244/62.
DR N-PSDB; AAA96769.
XX
PT Diagnosing an iron disorder e.g. hemochromatosis or a genetic
PT susceptibility to develop it, by determining the presence of a mutation
PT in exon 2 or an intron of a histocompatibility iron loading nucleic
PT acid -
XX
PS Disclosure; Page 3; 55pp; English.
XX
CC The present sequence represents a human histocompatibility iron loading
CC (HFE) protein. The HFE gene is a major histocompatibility (MHC)
CC non-classical class I gene located on chromosome 6p. Mutations in the
CC gene lead to iron disorders. The specification describes a method for
CC diagnosing an iron disorder or a genetic susceptibility to develop the
CC disorder in a mammal. The method comprises determining the presence of
CC a mutation in exon 2 or an intron of a HFE gene or protein. The mutation
CC is not a C to G missense mutation at nucleotide 187 of the sequence
CC given in A96769 (Genbank Accession number U60319). The presence of the
CC mutation indicates the disorder or the genetic susceptibility to the
CC disorder. The method is used to diagnose an iron disorder
CC e.g. haemochromatosis, or a genetic susceptibility to develop it.
XX
SQ Sequence 348 AA;
Query Match 99.5%; Score 1513; DB 21; Length 348;
Best Local Similarity 99.6%; Pred. No. 4.5e-134;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISSQ 82
QY 61 MWLQSLQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCMQEDNSTEGYWKYGYDG 142
QY 121 QDHLEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNAYLERDCAQQLLELGRGVL 180
DB 143 QDHLEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNAYLERDCAQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSSVTLRCLALNYPQNTMKWLKDKQPMDAKEPEKDPVLPNGDG 240
DB 203 DQOVPLVKVTHVTSSVTLRCLALNYPQNTMKWLKDKQPMDAKEPEKDPVLPNGDG 262
QY 241 TYQGWTITLAVPPGEGORYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWTITLAVPPGEGORYTCQVEHPGLDQPLIVWE 298
RESULT 8
AAB36869

ID AAB36869 standard; Protein; 348 AA.
XX
AC AAB36869;
XX
DT 21-FEB-2001 (first entry)
XX
DE Human hereditary hemochromatosis protein.
XX
KW HH; hereditary hemochromatosis; chelation agent;
KW T-cell differentiation factor; iron overload.
XX
OS Homo sapiens.
XX
PN US6140305-A.
XX
PD 31-OCT-2000.
XX
PF 04-APR-1997; 97US-0834497.
XX
PR 04-APR-1996; 96US-0630912.
PR 16-APR-1996; 96US-0632673.
PR 23-MAY-1996; 96US-0652265.
XX
PA (BIRA) BIO-RAD LAB INC.
XX
PI Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
PI Feder JN;
XX
DR WPI; 2001-006341/01.
DR N-PSDB; AAC68425.
XX
PT New hereditary hemochromatosis gene products or polypeptides, useful
PT for treating hereditary hemochromatosis in a patient, and as a metal
PT chelation agent alleviating iron overload -
XX
XX
PS Claim 1; Fig 4; 108pp; English.
XX
CC The present invention relates to hereditary hemochromatosis gene
CC products. These proteins may be used to treat a patient diagnosed as
CC having human hemochromatosis disease. It is also useful as a metal
CC chelation agent or as a T-cell differentiation factor, and for
CC alleviating iron overload. They may also be used in protein replacement
CC therapy for individuals having a defective human hemochromatosis gene.
XX
SQ Sequence 348 AA;
Query Match 99.5%; Score 1513; DB 22; Length 348;
Best Local Similarity 99.6%; Pred. No. 4.5e-134;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISQ 60
DB 23 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISQ 82
QY 61 MWLQLSQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCFDPDLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCAQQLLELGRGVL 180
DB 143 QDHLFCFDPDLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCAQQLLELGRGVL 202
QY 181 DQOVPLPVKVTHTVSSVTLTLCRALNYYPQNTMKWLKDQPMDAKEFEPKDVLPNGDG 240
DB 203 DQOVPLPVKVTHTVSSVTLTLCRALNYYPQNTMKWLKDQPMDAKEFEPKDVLPNGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 9
AAB36872

ID AAB36872 standard; Protein; 348 AA.
XX
AC AAB36872;
XX
DT 21-FEB-2001 (first entry)
XX
DE Human hereditary hemochromatosis 24d1/2 mutation protein.
XX
KW HH; hereditary hemochromatosis; chelation agent;
KW T-cell differentiation factor; iron overload.
XX
OS Homo sapiens.
XX
PN US6140305-A.
XX
PD 31-OCT-2000.
XX
PF 04-APR-1997; 97US-0834497.
XX
PR 04-APR-1996; 96US-0630912.
PR 16-APR-1996; 96US-0632673.
PR 23-MAY-1996; 96US-0652265.
XX
PA (BIRA) BIO-RAD LAB INC.
XX
PI Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
PI Feder JN;
XX
DR WPI; 2001-006341/01.
DR N-PSDB; AAC68428.
XX
PT New hereditary hemochromatosis gene products or polypeptides, useful
PT for treating hereditary hemochromatosis in a patient, and as a metal
PT chelation agent alleviating iron overload -
XX
XX
PS Claim 4; Fig 4; 108pp; English.
XX
CC The present invention relates to hereditary hemochromatosis gene
CC products. These proteins may be used to treat a patient diagnosed as
CC having human hemochromatosis disease. It is also useful as a metal
CC chelation agent or as a T-cell differentiation factor, and for
CC alleviating iron overload. They may also be used in protein replacement
CC therapy for individuals having a defective human hemochromatosis gene.
XX
SQ Sequence 348 AA;
Query Match 99.3%; Score 1509; DB 22; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.1e-133;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISQ 60
DB 23 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISQ 82
QY 61 MWLQLSQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCFDPDLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCAQQLLELGRGVL 180
DB 143 QDHLFCFDPDLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCAQQLLELGRGVL 202
QY 181 DQOVPLPVKVTHTVSSVTLTLCRALNYYPQNTMKWLKDQPMDAKEFEPKDVLPNGDG 240
DB 203 DQOVPLPVKVTHTVSSVTLTLCRALNYYPQNTMKWLKDQPMDAKEFEPKDVLPNGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 10
AAB36872

ID AC AAU80035 standard; Protein; 438 AA.
XX AAU80035;
DT 15-JUL-2002 (first entry)
XX Beta 2 microglobulin (beta2M)/HFE monochain.
DE Human; beta 2 microglobulin; beta2M/HFE monochain; HFE; ischaemia;
XX iron absorption regulator; intracellular iron absorption; lung injury;
KW haemochromatosis; transfusion; thalassaemia; haemolytic anaemia;
KW chronic infection; transferrin receptor; Tfr; brain tumour; cancer;
KW oxidative stress disorder; tissue damage; vascular disease;
KW inflammation; atherosclerosis; autoimmune disease;
XX inflammatory condition.
XX Homo sapiens.
OS WO200224929-A2.
XX 28-MAR-2002.
XX 24-SEP-2001; 2001WO-US29873.
XX 22-SEP-2000; 2000US-234843P.
XX (UYRA-) UNIV RAMOT APPLIED RES & IND DEV LTD.
PA (MCIN/) MCINNIS P.
XX Ehrlich R, Rotem-Yehudar R, Laham N;
PI WPI; 2002-383192/41.
XX N-PSDB; ABK49917.
DR Soluble beta 2 microglobulin/HFE monochain useful for treating
PT iron-overload conditions e.g. thalassaemia and chronic infections,
PT comprises human beta 2 microglobulin linked to alpha domains of HFE by
PT a linker peptide -
XX Example 2; Fig 2; 77pp; English.
XX The invention relates to a soluble polypeptide (I) of beta 2
CC microglobulin (beta2M)/HFE monochain comprising human beta2M (or its
CC analogue or active fragment), linked to alpha1-alpha3 domains of human
CC HFE (a central regulator of iron absorption; undefined), or its analogue
CC or active fragment, by a flexible linker peptide, or a functional
CC derivative or salt of (I). (I) is useful for reducing intracellular iron
CC absorption in patients having hereditary haemochromatosis, transfusions,
CC thalassaemias, haemolytic anaemia or chronic infections, and for
CC delivering a therapeutic to cells that over-express transferrin receptor
CC (Tfr) which are preferably lymphocytes or leukocytes, across the blood-
CC brain barrier. (I) is further useful for treating brain tumour. (I)
CC is also useful for treating oxidative stress disorders resulting in
CC tissue damage e.g. vascular diseases, inflammation, atherosclerosis,
CC lung injury, ischaemia, etc. A DNA molecule (II) encoding (I) is useful
CC as a platform for drug delivery of therapeutic use for cancer,
CC autoimmune diseases and inflammatory conditions. The monochain manifests
CC specific characteristics advantageous for drug delivery systems. It is a
CC soluble, stable and fully conformed protein. It binds specifically to
CC transferrin receptor (Tfr) and therefore targets cells that over-express
CC this receptor. It is continuously internalised by the target cells, thus
CC enabling efficient drug delivery. It dissociates from the receptor in the
CC cells, minimising side effects. It negatively regulates iron absorption,
CC reducing growth of undesired cells and preventing lymphocyte activation.
CC It is not diluted in the blood as is transferrin. It should not induce an
CC immune response since it is a self non-polymorphic protein and delivery of
CC drugs via monochain is expected to overcome drug-resistance since it is a
CC natural Tfr-binding protein. The present sequence represents the amino
CC acid sequence of beta2M/HFE monochain.
XX Sequence 438 AA;
SQ

Query Match 99.2%; Score 1508; DB 23; Length 438;

Best Local Similarity 99.6%; Pred. No. 1.8e-133;
Matches 274; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLIRSHSLHYLFMGASEQDLGLSLFALGYVDDQLFVFFYDDERRRVERPTPWSSRISSQ 60
DB 135 RLIRSHSLHYLFMGASEQDLGLSLFALGYVDDQLFVFFYDDERRRVERPTPWSSRISSQ 194
QY 61 MWLQLSQSLKGMDFMTVDFTWIMENHNHKSHTTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 195 MWLQLSQSLKGMDFMTVDFTWIMENHNHKSHTTLQVILGCEMQEDNSTEGYWKYGYDG 254
QY 121 QHLEFCPTDLWRAAEPRAPWTKLEWERHKTARONRAYLERDCPAQLOQLLELGRGVL 180
DB 255 QHLEFCPTDLWRAAEPRAPWTKLEWERHKTARONRAYLERDCPAQLOQLLELGRGVL 314
QY 181 DQOVPLVKVTHHTVSSVTLRCALNYPQNTMKWLKDKQPMDAKEPEKDVLPNGDG 240
DB 315 DQOVPLVKVTHHTVSSVTLRCALNYPQNTMKWLKDKQPMDAKEPEKDVLPNGDG 374
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIM 275
DB 375 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIM 409
RESULT 11
AAB36870
ID AAB36870 standard; Protein; 348 AA.
XX AAB36870;
AC AAB36870;
XX 21-FEB-2001 (first entry)
DT Human hereditary hemochromatosis 24d1 mutation protein.
DE HH; hereditary hemochromatosis; chelation agent;
KW T-cell differentiation factor; iron overload.
XX Homo sapiens.
OS US6140305-A.
PN 31-OCT-2000.
PD 04-APR-1997; 97US-0834497.
XX 04-APR-1996; 96US-0630912.
PR 16-APR-1996; 96US-0632673.
PR 23-MAY-1996; 96US-0652265.
XX (BIRA) BIO-RAD LAB INC.
PA Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
PI Feder JN;
XX WPI; 2001-006341/01.
DR N-PSDB; AAC68426.
XX New hereditary hemochromatosis gene products or polypeptides, useful
PT for treating hereditary hemochromatosis in a patient, and as a metal
PT chelation agent alleviating iron overload -
XX Claim 2; Fig 3; 108pp; English.
XX The present invention relates to hereditary hemochromatosis gene
CC products. These proteins may be used to treat a patient diagnosed as
CC having human hemochromatosis disease. It is also useful as a metal
CC chelation agent or as a T-cell differentiation factor, and for
CC alleviating iron overload. They may also be used in protein replacement
CC therapy for individuals having a defective human hemochromatosis gene.
XX Sequence 348 AA;
SQ

Query Match 98.8%; Score 1502; DB 22; Length 348;

Best Local Similarity 99.3%; Pred. No. 4.9e-133;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLPMGASQDGLSLFEALGYDDQLFVYDDESRVPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLPMGASQDGLSLFEALGYDDQLFVYDDESRVPRTPWVSSRISSQ 82
QY 61 MWLQLSQSLKGWDMFTVDFWTIMENHNHKSHTLQVILGCEMOEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGWDMFTVDFWTIMENHNHKSHTLQVILGCEMOEDNSTEGYWKYGYDG 142
QY 121 QDHLFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDPCPAQLQQLLELGRGVL 180
DB 143 QDHLFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDPCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHHTVSSVTTLRCRALNYPQNTMKWLKDKQPMADAKEFPKDVLPNGDG 240
DB 203 DQVPPPLVKVTHHTVSSVTTLRCRALNYPQNTMKWLKDKQPMADAKEFPKDVLPNGDG 262
QY 241 TYQGWITLAVPPGEGEORYTCQVEHPGLDQPLIVIE 276
DB 263 TYQGWITLAVPPGEGEORYTCQVEHPGLDQPLIVIE 298

RESULT 12
AAW94297
ID AAW94297 standard; peptide; 276 AA.
XX AAW94297;
XX
XX
XX 27-APR-1999 (first entry)
XX
DE HFE mutant (H111A/H145A-HFE) polypeptide sequence.
XX
KW HFE; beta-2-microglobulin; beta2m; iron overload; haemochromatosis;
KW transfusion; protein replacement therapy; hereditary haemochromatosis;
KW transferrin receptor; iron deficiency; anemia; mutant.
XX
OS Synthetic.
XX
XX
FH Key Location/Qualifiers
FT Misc-difference 2 /note= "indicated in the sequence listing as Arg"
FT
FT Misc-difference 89 /label= H111A
FT /note= "wild type His (of the mature protein sequence)
FT /note= "is replaced by Ala"
FT Misc-difference 123 /label= H145A
FT /note= "wild type His (of the mature protein sequence)
FT /note= "is replaced by Ala"
XX
XX WO9856814-A1.
XX
XX
XX 17-DEC-1998.
XX
XX 12-JUN-1998; 98WO-US12436.
XX
XX 13-JUN-1997; 97US-0876010.
XX
XX (CALY) CALIFORNIA INST OF TECHNOLOGY.
XX (PROG-) PROGENITOR INC.
XX
XX Bjorkman PJ, Feder JN, Schatzman RC;
XX
XX WPI; 1999-080886/07.
XX
XX New treatment of an iron overload disease - comprises use of HFE
XX polypeptides provided in a complex with full length, wild type human
XX (2m), useful in protein replacement therapy
XX
XX Claim 5; Page 15; 36pp; English.

CC The present sequence represents a H111A/H145A-HFE mutant polypeptide.
CC The HFE polypeptides (AAW94295-297) provided in a complex with full
CC length, wild type human beta-2-microglobulin (beta2m) form compositions
CC in the treatment of primary iron overload diseases (e.g.
CC haemochromatosis), or other iron overload conditions resulting from
CC secondary causes (e.g. repeated transfusions). Data regarding the
CC structure and function correlations of HFE polypeptides is useful in
CC designing drugs that modulate the HFE gene and HFE activity. The
CC polypeptides are also useful in protein replacement therapy for
CC individuals possessing a defective HFE gene (e.g. Hereditary
CC haemochromatosis). (Ant)agonists of the polypeptides are also useful in
CC treating primary and secondary iron overload diseases. The modulators of
CC the transferrin receptor are useful in treating iron deficiency
CC conditions such as anemia, and in modulating the amount of iron
CC transported into a cell. The HFE polypeptides provide a molecular basis
CC for the relationship between HFE and iron metabolism, which enables
CC treatment of iron overload and deficiency diseases.
XX
SQ Sequence 276 AA;

Query Match 98.2%; Score 1493; DB 20; Length 276;
Best Local Similarity 98.9%; Pred. No. 2.5e-132;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLPMGASQDGLSLFEALGYDDQLFVYDDESRVPRTPWVSSRISSQ 60
DB 1 RLLRSHSLHYLPMGASQDGLSLFEALGYDDQLFVYDDESRVPRTPWVSSRISSQ 60
QY 61 MWLQLSQSLKGWDMFTVDFWTIMENHNHKSHTLQVILGCEMOEDNSTEGYWKYGYDG 120
DB 61 MWLQLSQSLKGWDMFTVDFWTIMENHNHKSHTLQVILGCEMOEDNSTEGYWKYGYDG 120
QY 121 QDHLFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDPCPAQLQQLLELGRGVL 180
DB 121 QDHLFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDPCPAQLQQLLELGRGVL 180
QY 181 DQVPPPLVKVTHHTVSSVTTLRCRALNYPQNTMKWLKDKQPMADAKEFPKDVLPNGDG 240
DB 181 DQVPPPLVKVTHHTVSSVTTLRCRALNYPQNTMKWLKDKQPMADAKEFPKDVLPNGDG 240
QY 241 TYQGWITLAVPPGEGEORYTCQVEHPGLDQPLIVIE 276
DB 241 TYQGWITLAVPPGEGEORYTCQVEHPGLDQPLIVIE 276

RESULT 13
ABG72687
ID ABG72687 standard; protein; 276 AA.
XX
XX ABG72687;
XX
XX 05-MAR-2003 (first entry)
XX
XX Human haemochromatosis (HFE) mature protein, mutant H89A/H123A.
XX
XX Human; haemochromatosis; HFE; hereditary haemochromatosis;
KW iron overload disease; iron deficiency disease; Beta2-microglobulin;
KW Beta2m; transferrin receptor; anaemia; mutant; muten.
XX
XX Homo sapiens.
OS Synthetic.
XX
XX Key Location/Qualifiers
FH Misc-difference 89 /note= "wild-type His substituted by Ala"
FT Misc-difference 123 /note= "wild-type His substituted by Ala"
FT
XX US6391852-B1.
XX
XX 21-MAY-2002.
XX
XX 12-JUN-1998; 98US-0094964.

XX PR 13-JUN-1997; 97US-0876010.
XX PA (BIRA) BIO-RAD LAB INC.
XX PA (CALY) CALIFORNIA INST OF TECHNOLOGY.
XX PI Feder JN, Bjorkman PJ, Schatzman RC;
XX WPI; 2003-155377/15.
XX
XX Method of treating an iron overload disease comprises administration of
XX a soluble complex comprising a 276 amino acid HFE polypeptide and a
XX full length, wild-type human beta2m -
XX
XX Claim 3; Column 2; 17pp; English.
XX
XX The invention relates to a method of treating an iron overload disease
XX comprising administration of a soluble complex comprising a 276 amino
XX acid mature HFE (hereditary haemochromatosis gene protein) polypeptide
XX (ABG72685-ABG72687) and a full length, wild-type human beta2m
XX (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of
XX ⁵¹Fe-transferrin in the presence of purified H63D-HFE/beta2m
XX heterodimers was determined. At a concentration of 250 nM H63D-HFE/
XX beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for
XX transferrin of 28 nM. At the same concentration of normal HFE/beta2m
XX heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence
XX of any HFE/beta2m heterodimers, TfR displayed a KD for transferrin of
XX 7nM. It was observed that H63D-HFE/beta2m heterodimers were 30-40 % less
XX efficient in decreasing TfR affinity for transferrin compared to
XX wild-type HFE. The method is useful for treating iron overload diseases
XX and iron deficiency e.g. anaemia. The present sequence is the H11A/H145A
XX (residues 111 and 145 of the full length protein, 89/123 of the mature
XX form) mutant from of mature HFE used to investigate the role of the His
XX residues in transferrin receptor binding to transferrin.
XX
XX SQ Sequence 276 AA;

Query Match 98.2%; Score 1493; DB 24; Length 276;
Best Local Similarity 98.9%; Pred. No. 2.5e-132;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSLHLYFMGASEODLGLSLFEALGYVDQLFVYDDESRVPRTPWSSRISSQ 60
DB 1 RLLRSLHLYFMGASEODLGLSLFEALGYVDQLFVYDDESRVPRTPWSSRISSQ 60
QY 61 MWLQSLKGDHMFVDFWTFIMENHNSKESHILQVILGCEMOEDNSTEGYWKYGDG 120
DB 61 MWLQSLKGDHMFVDFWTFIMENHNSKESHILQVILGCEMOEDNSTEGYWKYGDG 120
QY 121 QDHFECPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 121 QDALEFCPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQOVPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEFEFPKDVLPNGDG 240
DB 181 DQOVPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEFEFPKDVLPNGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVHPGLDQPLIWIWE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVHPGLDQPLIWIWE 276

RESULT 14
AAB36873
ID AAB36873 standard; Protein; 361 AA.
XX
XX AAB36873;
XX
XX 21-FEB-2001 (first entry)
XX
XX Rabbit leukocyte antigen.
XX
XX HH; hereditary hemochromatosis; chelation agent;
KW

KW T-cell differentiation factor; iron overload.
XX
XX Oryctolagus cuniculus.
XX
XX US6140305-A.
XX 31-OCT-2000.
XX
XX 04-APR-1997; 97US-08344497.
XX
XX 04-APR-1996; 96US-0630912.
XX 16-APR-1996; 96US-0632873.
XX 23-MAY-1996; 96US-0652265.
XX
XX (BIRA) BIO-RAD LAB INC.
XX
XX Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
XX Feder JN;
XX WPI; 2001-006341/01.
XX
XX New hereditary hemochromatosis gene products or polypeptides, useful
XX for treating hereditary hemochromatosis in a patient, and as a metal
XX chelation agent alleviating iron overload -
XX
XX Disclosure; Fig 7; 108pp; English.
XX
XX The present invention relates to hereditary hemochromatosis gene
XX products. These proteins may be used to treat a patient diagnosed as
XX having human hemochromatosis disease. It is also useful as a metal
XX chelation agent or as a T-cell differentiation factor, and for
XX alleviating iron overload. They may also be used in protein replacement
XX therapy for individuals having a defective human hemochromatosis gene.
XX
XX SQ Sequence 361 AA;
Query Match 34.4%; Score 523; DB 22; Length 361;
Best Local Similarity 40.1%; Pred. No. 9.1e-41;
Matches 111; Conservative 44; Mismatches 114; Indels 8; Gaps 7;
QY 5 SHSLHLYFMGASEODLGLSLFEALGYVDQLFVYDDE--SRVPRTPWSSRISSQMW 62
DB 26 SHSMRYFTYSVRPGELGEPRTIIVGYVDPTQVRFDSDAASPRMEQRAFM--GQVEPEY 84
QY 63 LQLSLSKGDHMFVDFWTFIMENHNSKE--SHILQVILGCEMOEDNS--TEGYWKYGDG 120
DB 85 DQQTQIAKDTAQTFRVNLNTALRYNQSAAGSHFTFTMGCEVWADGRFFHGYRAYDG 144
QY 121 QDHFECPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 ADVIALNEDLRSWTAADTAQNTQKWEAAG--EAEHRHAYLERECVEWLRRLVLEMGKETL 203
QY 181 DQOVPLVKVTHVTSS--VTLRCALNYPQNTMKWLKDKQPMDAKEFEFPKDVLPNGD 239
DB 204 QRADPPKAVHTHPASDREATLRCAWLGFPYPAISLTWQDGDG--QTQDTTELVTETPGD 262
QY 240 GTYQGWITLAVPPGEEQRYTCQVHPGLDQPLIWIWE 276
DB 263 GTYQKAAVVPVSGEQRVTCVQHEGLPEPLTLTWE 299
RESULT 15
ABP68379
ID ABP68379 standard; Protein; 92 AA.
XX
XX ABP68379;
XX
XX 08-JAN-2003 (first entry)
XX
XX Human colon specific protein, SEQ ID 120.
XX
XX Human; colon; cytostatic; vaccine; colon cancer; colon disorder;
KW metastasis.

[illegible]

Search completed: August 5, 2003, 13:08:23
Job time : 39 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:07:04 ; Search time 14.5 seconds
(without alignments)
805.365 Million cell updates/sec

Title: US-10-092-404-2
Perfect score: 1520
Sequence: 1 RLLRSHLHFLWFGASEQDL.....RYTCQVEHFGDQPLIVWE 276

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 42310858 residues

Total number of hits satisfying chosen parameters: 328717

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents AA:*
1: /cgn2_6/ptodata/1/iaa/5A COMB.pap.*
2: /cgn2_6/ptodata/1/iaa/5B COMB.pap.*
3: /cgn2_6/ptodata/1/iaa/6A COMB.pap.*
4: /cgn2_6/ptodata/1/iaa/6B COMB.pap.*
5: /cgn2_6/ptodata/1/iaa/PCTUS COMB.pap.*
6: /cgn2_6/ptodata/1/iaa/backfile1.pap.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1520	100.0	276	4	US-09-094-964-2 Sequence 2, Appli
2	1520	100.0	348	3	US-08-552-265-6 Sequence 6, Appli
3	1520	100.0	348	3	US-08-834-497A-6 Sequence 6, Appli
4	1520	100.0	348	3	US-09-503-444A-6 Sequence 1, Appli
5	1513	99.5	276	4	US-09-094-964-1 Sequence 1, Appli
6	1513	99.5	348	3	US-08-552-265-2 Sequence 2, Appli
7	1513	99.5	348	3	US-08-834-497A-2 Sequence 2, Appli
8	1513	99.5	348	3	US-09-503-444A-2 Sequence 2, Appli
9	1513	99.5	348	4	US-09-277-457-2 Sequence 2, Appli
10	1513	99.5	348	4	US-09-679-729-2 Sequence 2, Appli
11	1509	99.3	348	3	US-08-552-265-8 Sequence 8, Appli
12	1509	99.3	348	3	US-08-834-497A-8 Sequence 8, Appli
13	1509	99.3	348	3	US-09-503-444A-8 Sequence 8, Appli
14	1502	98.8	348	3	US-08-552-265-4 Sequence 4, Appli
15	1502	98.8	348	3	US-08-834-497A-4 Sequence 4, Appli
16	1502	98.8	348	3	US-09-503-444A-4 Sequence 4, Appli
17	1493	98.2	276	4	US-09-094-964-3 Sequence 3, Appli
18	523	34.4	361	3	US-08-552-265-22 Sequence 22, Appl
19	523	34.4	361	3	US-08-834-497A-22 Sequence 22, Appl
20	523	34.4	361	3	US-09-503-444A-22 Sequence 22, Appl
21	517	34.0	364	4	US-08-514-372C-11 Sequence 11, Appl
22	514	33.8	365	3	US-08-552-265-23 Sequence 23, Appl
23	514	33.8	365	3	US-08-834-497A-23 Sequence 23, Appl
24	514	33.8	365	3	US-09-503-444A-23 Sequence 23, Appl
25	506	33.3	274	2	US-08-484-905-107 Sequence 107, App
26	506	33.3	274	3	US-08-481-985B-107 Sequence 107, App
27	506	33.3	274	3	US-08-370-476-107 Sequence 107, App

28	506	33.3	341	3	US-08-890-719-38 Sequence 38, Appl
29	505	33.2	365	2	US-08-484-905-97 Sequence 97, Appl
30	505	33.2	365	3	US-08-481-985B-97 Sequence 97, Appl
31	505	33.2	365	3	US-08-370-476-97 Sequence 97, Appl
32	504	33.2	274	2	US-08-484-905-108 Sequence 108, App
33	504	33.2	274	3	US-08-481-985B-108 Sequence 108, App
34	504	33.2	274	3	US-08-370-476-108 Sequence 108, App
35	504	33.2	365	2	US-08-484-905-100 Sequence 100, App
36	504	33.2	365	3	US-08-481-985B-100 Sequence 100, App
37	504	33.2	365	3	US-08-370-476-100 Sequence 100, App
38	503	33.1	274	1	US-08-222-851-1 Sequence 1, Appli
39	503	33.1	363	4	US-08-914-372C-37 Sequence 37, Appl
40	503	33.1	365	2	US-08-484-905-99 Sequence 99, Appl
41	503	33.1	365	3	US-08-481-985B-99 Sequence 99, Appl
42	503	33.1	365	3	US-08-370-476-99 Sequence 99, Appl
43	502	33.0	274	2	US-08-484-905-106 Sequence 106, App
44	502	33.0	274	3	US-08-481-985B-106 Sequence 106, App
45	502	33.0	274	3	US-08-370-476-106 Sequence 106, App

ALIGNMENTS

RESULT 1
US-09-094-964-2
; Sequence 2, Application US/09094964
; Patent No. 6391852
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; APPLICANT: Bjorkman, Pamela J.
; APPLICANT: Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; TITLE OF INVENTION: AND IRON DEFICIENCY DISEASES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSES: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-09-094-964-2

Query Match 100.0%; Score 1520; DB 4; Length 276;
Best Local Similarity 100.0%; Pred. No. 2.2e-142;

Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWSSRISSQ 60
DB 1 RLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWSSRISSQ 60

QY 61 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120

QY 121 QDLHFPCDPTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 121 QDLHFPCDPTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180

QY 181 DQOVPLVAVKTHVTSSVTLRCALNYPQNTMKWLKDKQMDAKEPEPKDVLPGDG 240
DB 181 DQOVPLVAVKTHVTSSVTLRCALNYPQNTMKWLKDKQMDAKEPEPKDVLPGDG 240

QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 2

US-08-652-265-6
; Sequence 6, Application US/08652265
; Patent No. 6025130
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M.
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 17957-000500
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-652-265-6

Query Match 100.0%; Score 1520; DB 3; Length 348;
Best Local Similarity 100.0%; Pred. No. 3e-142;
Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWSSRISSQ 60

DB 23 RLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWSSRISSQ 82
QY 61 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142

QY 121 QDLHFPCDPTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDLHFPCDPTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202

QY 181 DQOVPLVAVKTHVTSSVTLRCALNYPQNTMKWLKDKQMDAKEPEPKDVLPGDG 240
DB 203 DQOVPLVAVKTHVTSSVTLRCALNYPQNTMKWLKDKQMDAKEPEPKDVLPGDG 262

QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 3

US-08-834-497A-6
; Sequence 6, Application US/08834497A
; Patent No. 6149305
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
; NUMBER OF SEQUENCES: 76
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/834,497A
; FILING DATE: 04-APR-1997
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/632,673
; FILING DATE: 16-APR-1996
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/630,912
; FILING DATE: 04-APR-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0056-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids


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;
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-834-497A-6

Query Match 100.0%; Score 1520; DB 3; Length 348;
Best Local Similarity 100.0%; Pred. No. 3e-142;
Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDLGLSLFEALGYVDQDLFFVYDDERRRVEPRTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASQDLGLSLFEALGYVDQDLFFVYDDERRRVEPRTPWSSRISSQ 82
QY 61 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMDEDSSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMDEDSSTEGYWKYGYDG 142
QY 121 QDHLEFCPTDLWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPTDLWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 203 DQOVPLVKVTHVTSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 298

RESULT 4
US-09-503-444A-6
; Sequence 6, Application US/09503444A
; Patent No. 6228594
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: Wordperfect Version 8
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/503,444A
; FILING DATE: 14-Feb-2000
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/652,265
; FILING DATE: 23-May-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,673
; FILING DATE: 16-Apr-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/630,912
; FILING DATE: 04-Apr-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0088-999
```

```
;
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 212-790-9090
; TELEFAX: 212-869-9741
; TELEX: 66141
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-09-503-444A-6

Query Match 100.0%; Score 1520; DB 3; Length 348;
Best Local Similarity 100.0%; Pred. No. 3e-142;
Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDLGLSLFEALGYVDQDLFFVYDDERRRVEPRTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASQDLGLSLFEALGYVDQDLFFVYDDERRRVEPRTPWSSRISSQ 82
QY 61 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMDEDSSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMDEDSSTEGYWKYGYDG 142
QY 121 QDHLEFCPTDLWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPTDLWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 203 DQOVPLVKVTHVTSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 298

RESULT 5
US-09-094-964-1
; Sequence 1, Application US/09094964
; Patent No. 6391852
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; APPLICANT: Bjorkman, Pamela J.
; APPLICANT: Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; NUMBER OF INVENTION: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:
```

TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-094-964-1

Query Match 99.5%; Score 1513; DB 4; Length 276;
Best Local Similarity 99.6%; Pred. No. 1.1e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQLFVYDDESRVPRTPWSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQLFVYDDESRVPRTPWSSRISSQ 60
QY 61 MWLQSLQSLKGDHMTVDFTIMENHNHSHKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
DB 61 MWLQSLQSLKGDHMTVDFTIMENHNHSHKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
QY 121 QDHLFCFDPDLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 121 QDHLFCFDPDLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 181 DQVPPPLVKVTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDPLIWIWE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDPLIWIWE 276

RESULT 6
US-08-652-265-2
Sequence 2, Application US/08652265
Patent No. 6025130
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Hereditary Hemochromatosis Gene
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend and Crew LLP
STREET: Two Embarcadero Center, Eighth Floor
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94111-3834
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Smith, William M.
REGISTRATION NUMBER: 30,223
REFERENCE/DOCKET NUMBER: 17957-000500
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 576-0200

TELEFAX: (415) 576-0300
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-652-265-2

Query Match 99.5%; Score 1513; DB 3; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.1e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQLFVYDDESRVPRTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQLFVYDDESRVPRTPWSSRISSQ 82
QY 61 MWLQSLQSLKGDHMTVDFTIMENHNHSHKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGDHMTVDFTIMENHNHSHKESHTLQVILGCEMOEDNSTEGYWKYGYDG 142
QY 121 QDHLFCFDPDLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLFCFDPDLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 203 DQVPPPLVKVTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDPLIWIWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDPLIWIWE 298

RESULT 7
US-08-834-497A-2
Sequence 2, Application US/08834497A
Patent No. 6140305
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FastSeq for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/834,497A
FILING DATE: 04-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/632,673
FILING DATE: 16-APR-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:

APPLICATION NUMBER: US 08/630,912
FILING DATE: 04-APR-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0056-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-834-497A-2

Query Match 99.5%; Score 1513; DB 3; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.5e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 82
QY 61 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QHLEFCPTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QHLEFCPTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPEPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 203 DQVPEPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPGEQRVTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPGEQRVTCQVEHPGLDQPLIVWE 298

RESULT 8
US-09-503-444A-2
Sequence 2, Application US/09503444A
Patent No. 6228594
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Hereditary Hemochromatosis Gene
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: WordPerfect Version 8
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/503,444A
FILING DATE: 14-Feb-2000

CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/652,265
FILING DATE: 23-May-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/632,673
FILING DATE: 16-Apr-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/630,912
FILING DATE: 04-Apr-1996
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0088-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-790-9090
TELEFAX: 212-869-9741
TELEX: 66141
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-503-444A-2

Query Match 99.5%; Score 1513; DB 3; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.5e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 82
QY 61 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QHLEFCPTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QHLEFCPTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPEPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 203 DQVPEPLVKVTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPGEQRVTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPGEQRVTCQVEHPGLDQPLIVWE 298

RESULT 9
US-09-277-457-2
Sequence 2, Application US/09277457
Patent No. 6355425
GENERAL INFORMATION:
APPLICANT: Rothenberg, Barry E.
APPLICANT: Sawada-Hirai, Ritsuko
APPLICANT: Barton, James C.
TITLE OF INVENTION: MUTATIONS ASSOCIATED WITH IRON DISORDERS
FILE REFERENCE: 10653/002001
CURRENT APPLICATION NUMBER: US/09/277,457
CURRENT FILING DATE: 1999-03-26
NUMBER OF SEQ ID NOS: 30
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO: 2
LENGTH: 348
TYPE: PRT
ORGANISM: Homo Sapiens
US-09-277-457-2
Query Match 99.5%; Score 1513; DB 4; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.5e-141;

Matches	275;	Conservative	0;	Mismatches	1;	Indels	0;	Gaps	0;
Qy	1	RLLRSHSLHYLFMGASEODLGLSLFEALGYVDDQLFVYDDSRVETPTPWSSRISSQ	60						
Db	23	RLLRSHSLHYLFMGASEODLGLSLFEALGYVDDQLFVYDDSRVETPTPWSSRISSQ	82						
Qy	61	MWQLQSLSKGDHMFVDFWTIMENHNHKSESHTLVILGCEMQEDNSTEGYKYGVDG	120						
Db	83	MWQLQSLSKGDHMFVDFWTIMENHNHKSESHTLVILGCEMQEDNSTEGYKYGVDG	142						
Qy	121	QDHLFPCPTLDWRAAPFPRAWPTKLEWERHKIRARONRAYLERDCAQLOOLLELGRGVL	180						
Db	143	QDHLFPCPTLDWRAAPFPRAWPTKLEWERHKIRARONRAYLERDCAQLOOLLELGRGVL	202						
Qy	181	DQQVPLVKVTHVHTSSVTTILRCALNYYPONITMKWLKDQPMDAKEFEKDVLPNGDG	240						
Db	203	DQQVPLVKVTHVHTSSVTTILRCALNYYPONITMKWLKDQPMDAKEFEKDVLPNGDG	262						
Qy	241	TYQGWITLAVPPGSEQRYTCQVHPGLDQPLIWIWE	276						
Db	263	TYQGWITLAVPPGSEQRYTCQVHPGLDQPLIWIWE	298						

```

RESULT 10
US-09-679-729-2
; Sequence 2, Application US/09679729
; Patent No. 6509442
; GENERAL INFORMATION:
; APPLICANT: Rothenberg, Barry E.
; APPLICANT: Sawada-Hirai, Ritsuko
; APPLICANT: Barton, James C.
; TITLE OF INVENTION: MUTATIONS ASSOCIATED WITH IRON DISORDERS
; FILE REFERENCE: 24065-004 DIV
; CURRENT APPLICATION NUMBER: US/09/679,729
; CURRENT FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: 09/277,457
; PRIOR FILING DATE: 1999-03-26
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 348
; TYPE: PRT
; ORGANISM: Homo Sapiens
US-09-679-729-2

```

[illegible]

RESULT 11
US-08-652-265-8
; Sequence 8, Application US/086522265

Patent No. 6025130

GENERAL INFORMATION:

APPLICANT: Thomas, Winston J.

APPLICANT: Drayna, Dennis T.

APPLICANT: Feder, John N.

APPLICANT: Gnirke, Andreas

APPLICANT: Ruddy, David

APPLICANT: Teuchinashi, Zenta

APPLICANT: Wolff, Roger K.

TITLE OF INVENTION: Hereditary Hemochromatosis Gene

NUMBER OF SEQUENCES: 44

CORRESPONDENCE ADDRESSES:

ADDRESSEE: Townsend and Townsend and Crew LLP

STREET: Two Embarcadero Center, Eighth Floor

CITY: San Francisco

STATE: California

COUNTRY: USA

ZIP: 94111-3834

COMPUTER READABLE FORM:

MEDIUM TYPE: Floppy disk

COMPUTER: IBM PC compatible

OPERATING SYSTEM: PC-DOS/MS-DOS

SOFTWARE: PatentIn Release #1.0, Version #1.30

CURRENT APPLICATION DATA:

APPLICATION NUMBER: US/08/652,265

FILING DATE: 23-MAY-1996

CLASSIFICATION: 514

ATTORNEY/AGENT INFORMATION:

NAME: Smith, William M.

REGISTRATION NUMBER: 30,223

REFERENCE/DOCKET NUMBER: 17957-000500

TELECOMMUNICATION INFORMATION:

TELEPHONE: (415) 576-0200

TELEFAX: (415) 576-0300

INFORMATION FOR SEQ ID NO: 8:

SEQUENCE CHARACTERISTICS:

LENGTH: 348 amino acids

TYPE: amino acid

TOPOLOGY: linear

MOLECULE TYPE: protein

US-0A-652-265-8

Query Match	99.3%;	Score 1509;	DB 3;	Length 348;
Best Local Similarity	99.6%;	Pred. No. 3.6e-141;		
Matches 275;	Conservative 0;	Mismatches 1;	Indels 0;	Gaps 0;
Qy	1	RLLRSHSLHYLFMGASEQDGLGLSIFEALGYVDQLFVYDDERRVDESRVPTPTWSSRISQ	60	
Db	23	RLLRSHSLHYLFMGASEQDGLGLSIFEALGYVDQLFVYDDERRVPTPTWSSRISQ	82	
Qy	61	MWLQLSLSKGDWHMFTVDFTWMENHNHKSESHLLQVILGCEMQEONSTEGYWKYGYDG	120	
Db	83	MWLQLSLSKGDWHMFTVDFTWMENHNHKSESHLLQVILGCEMQEONSTEGYWKYGYDG	142	
Qy	121	QDHLFCFCDTLDWRAEPRAWPTKLEWERHKIRARQNRAVYLERDCPAQLOQLLGLRGVL	180	
Db	143	QDHLFCFCDTLDWRAEPRAWPTKLEWERHKIRARQNRAVYLERDCPAQLOQLLGLRGVL	202	
Qy	181	DQQVPPPLVKVTHVTSSTVTLRCRALNYYPQNITMKWLKDQKPMDAKEFEPKQVLPNGDG	240	
Db	203	DQQVPPPLVKVTHVTSSTVTLRCRALNYYPQNITMKWLKDQKPMDAKEFEPKQVLPNGDG	262	
Qy	241	TYQGWITLAVPPGEEQRYTCQVHPGLDQPLIIVIE	276	
Db	263	TYQGWITLAVPPGEEQRYTCQVHPGLDQPLIIVIE	298	

RESULT 12
US-08-834-497A-8
; Sequence 8, Application US/08834497A
; Patent No. 6140305
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.

APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FASTSEQ for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/834,497A
FILING DATE: 04-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/632,673
FILING DATE: 16-APR-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/630,912
FILING DATE: 04-APR-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0056-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-834-497A-8

Query Match 99.3%; Score 1509; DB 3; Length 348;
Best Local Similarity 99.6%; Pred. No. 3.6e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLRLSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVETPTPWSSRISSQ 60
DB 23 RLRLSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVETPTPWSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMTFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCPTLDWRAAEPRAPWTKLEWERHKIRARQNAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLFCPTLDWRAAEPRAPWTKLEWERHKIRARQNAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQQVPLVKVTHVTSVTLRCAALNYPQNTMKWLKDKQPMDAKEPEPKDVLPNGDG 240
DB 203 DQQVPLVKVTHVTSVTLRCAALNYPQNTMKWLKDKQPMDAKEPEPKDVLPNGDG 262
QY 241 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIWIWE 276

DB 263 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIWIWE 298
RESULT 13
US-09-503-444A-8
Sequence 8, Application US/09503444A
Patent No. 6228594
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Hereditary Hemochromatosis Gene
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: WordPerfect Version 8
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/503,444A
FILING DATE: 14-Feb-2000
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/652,265
FILING DATE: 23-May-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/632,673
FILING DATE: 16-Apr-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/630,912
FILING DATE: 04-Apr-1996
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0088-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-790-9090
TELEFAX: 212-869-9741
TELEX: 66141
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-503-444A-8

Query Match 99.3%; Score 1509; DB 3; Length 348;
Best Local Similarity 99.6%; Pred. No. 3.6e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLRLSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVETPTPWSSRISSQ 60
DB 23 RLRLSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVETPTPWSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMTFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCPTLDWRAAEPRAPWTKLEWERHKIRARQNAYLERDCPAQLQQLLELGRGVL 180

Db 143 QDHLFCFCDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCAQLOQLLELGRGVL 202
Qy 181 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
Db 203 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
Qy 241 TYQGWITLAVPPGEEQRYTCQVEHGLDQPLIWIWE 276
Db 263 TYQGWITLAVPPGEEQRYTCQVEHGLDQPLIWIWE 298

RESULT 14
US-08-652-265-4
; Sequence 4, Application US/08652265
; Patent No. 6025130
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M.
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 17957-000500
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-652-265-4

Query Match 98.8%; Score 1502; DB 3; Length 348;
Best Local Similarity 99.3%; Pred. No. 1.8e-140;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 RLLRSHSLHYLPMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWVSSRISSQ 60
Db 23 RLLRSHSLHYLPMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWVSSRISSQ 82

Qy 61 MWLQSLKGLWDMFTVDFTWMENHNHSHKESHTLQVILGCMQEDNSTEGWVKYGYDG 120
Db 83 MWLQSLKGLWDMFTVDFTWMENHNHSHKESHTLQVILGCMQEDNSTEGWVKYGYDG 142

Qy 121 QDHLFCFCDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCAQLOQLLELGRGVL 180
Db 143 QDHLFCFCDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCAQLOQLLELGRGVL 202

Qy 181 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240

Db 203 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
Qy 241 TYQGWITLAVPPGEEQRYTCQVEHGLDQPLIWIWE 276
Db 263 TYQGWITLAVPPGEEQRYTCQVEHGLDQPLIWIWE 298

RESULT 15
US-08-834-497A-4
; Sequence 4, Application US/08834497A
; Patent No. 6140305
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
; NUMBER OF SEQUENCES: 76
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: FastSEQ for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/834,497A
; FILING DATE: 04-APR-1997
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/632,673
; FILING DATE: 16-APR-1996
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/630,912
; FILING DATE: 04-APR-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0056-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-834-497A-4

Query Match 98.8%; Score 1502; DB 3; Length 348;
Best Local Similarity 99.3%; Pred. No. 1.8e-140;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy 1 RLLRSHSLHYLPMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWVSSRISSQ 60
Db 23 RLLRSHSLHYLPMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWVSSRISSQ 82

Qy	61	MWLSQSLKGDHMFVDFWTFIMENHNHSHKESHTLQVILGCENQEDNSTEGYWKYCYDG	120
Db	83	MWLSQSLKGDHMFVDFWTFIMENHNHSHKESHTLQVILGCENQEDNSTEGYWKYCYDG	142
Qy	121	QDHLEFCPDTLDWEAEPRAWPTKLEWERHKIRARONRAYLERDCPAOLOLLELGRGVL	180
Db	143	QDHLEFCPDTLDWEAEPRAWPTKLEWERHKIRARONRAYLERDCPAOLOLLELGRGVL	202
Qy	181	DQVPPPLVKVTHVTSSVTTILRCALNYYQNITMKWLKDKQPMDAKEFEFEPKDVLPNGDG	240
Db	203	DQVPPPLVKVTHVTSSVTTILRCALNYYQNITMKWLKDKQPMDAKEFEFEPKDVLPNGDG	262
Qy	241	TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIWE	276
Db	263	TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIWE	298

Search completed: August 5, 2003, 13:11:11
Job time : 15.5 secs

